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(54) Title: GENOME OF *LEGIONELLA PNEUMOPHILA* PARIS AND LENS STRAIN-DIAGNOSTIC AND EPIDEMIOLOGICAL APPLICATIONS

(57) Abstract: The object of the invention is the genomic sequence and nucleotidic sequences coding for polypeptides of Legionella pneumophila Paris strain and Lens strain, such as cellular surface polypeptides, especially specific between these two strains and/or relative to the Philadelphia strain, or implied in the virulence or in the polysaccharide biosynthesis of cellular envelope, as well as vectors including said sequences and cells transformed by these vectors. The invention also concerns processes for detection of these nucleic acids or polypeptides and diagnostic typing kits for bacteria of the Legionella genre, especially of the Legionella pneumophila species, such as the Paris and Lens strains, between them and/or relative to the Philadelphia strain. The invention especially concerns a repeated nucleic sequence specific to the Legionella pneumophila species and its utilization as an analysis target in processes for detection of the presence of these bacteria. The aim of the invention is also a method for selection of compounds capable of modulating the biosynthesis of these polysaccharides of cellular envelope utilizing said nucleotidic sequences or said polypeptides. The invention finally comprises pharmaceutical compositions, especially vaccinal, for the prevention and/or treatment of bacterial infections, in particular by Legionella pneumophila Paris strain and/or Lens strain.



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GENOME OF *LEGIONELLA PNEUMOPHILA* PARIS AND LENS STRAIN - DIAGNOSTIC AND EPIDEMIOLOGICAL APPLICATIONS

The object of the invention is the genomic sequence and nucleotidic sequences coding for polypeptides of Legionella pneumophila Paris and Lens strain, such as polypeptides of cellular surface, especially specific between these two strains and/or relative to the Philadelphia strain, or implied in the virulence or in the biosynthesis of polysaccharides having a cellular envelope, as well as vectors including said sequences and cells transformed by these vectors. The invention likewise concerns processes for detection of these nucleic acids or polypeptides and diagnostic kits or kits for typing bacteria of the Legionella genre, especially the Legionella pneumophila species, such as the Paris and Lens strain, between them and/or relative to the Philadelphia strain. The invention especially concerns a specific repeated nucleic sequence of the Legionella pneumophila species and its utilization as an analysis target in processes for detection of the presence of these bacteria. The aim of the invention also is a method for selection of compounds capable of modulating the biosynthesis of these polysaccharides having a cellular envelope utilizing said nucleotidic sequences or said polypeptides. The invention finally comprises des pharmaceutical compositions, especially vaccinal, for the prevention and/or treatment of bacterial infections, in particular by the Legionella pneumophila Paris and/or Lens strain.

Legionella is a bacteria of the environment responsible for legionellosis and Pontiac fever. The epidemiological data indicate that probably only certain isolates are capable of causing clinical cases. The L. pneumophila species seems to have a more significant virulence than the other species, by being responsible for 90% of the cases of legionellosis. At the centre of this species, among the fifteen serogroups, the isolates of serogroup 1 are associated with 80% of cases.

To date, transmission from person to person has never been observed and measures for preventing legionellosis are thus concentrated on elimination of this pathogen from water circulation or from water-cooling towers in air-conditioning systems. In order to establish a rational policy for prevention, it is necessary to prevent the risk associated with each strain. In this optic, it would be desirable to be able to have processes or diagnostic kits available based on the recognition of protein or specific nucleic acid of this genre legio species pathogen or of a particular strain (or again subspecies) of this species.

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In effect, the interest in using these specific sequences in the domain of diagnostics or epidemiology rests on the possibility of analyzing a large number of sequences at the same time and very rapidly for:

- classification or typing of bacteria as a function of the presence of a sequence or of a profile of sequences characteristic of a genre, species or strain (sub-species) of bacteria, in particular in association with the gravity or not of pathologies which such bacteria can induce in case of infection in mammals, especially in humans; and

- simultaneous comparison of sequence or profile of sequences between different genres, species or strain (sub-species) of bacteria, pathogenic or not, especially enabling identification of a gene, or the corresponding proteic sequence, or a profile of genes whereof the presence and/or expression in a bacteria is specific to a genre, species or strain of bacteria, and/or to its pathogenicity or not, especially by means of tools such as DNA chips or, if required, protein chips, on which these specific sequences are immobilized.

These specific sequences can be specific sequences of the *Legionella* genre, or of a pathogenic bacteria of the *Legionella* genre and/or of the *Legionella pneumophila* species, or again of a bacteria of the *Legionella pneumophila* species Paris and/or Lens strain or again specific to a bacteria of the *Legionella pneumophila* species Paris and/or Lens strain relative to the Philadelphia strain.

This information is widely utilized especially to rapidly identify the presence or not of the pathogenic bacteria, the gravity of the infection which it can cause, the treatment adapted to an infection, and/or the necessity and the means to be put in place to decontaminate the objects, circuits or fluids which are contaminated or could be contaminated This information will likewise be widely used for epidemiological studies relative to this genre of bacteria.

This is just one of the aspects of the present invention.

In a first stage, the inventors have studied and attempted to comprehend the genomic diversity at the centre of the Legionella genre by complete sequencing of the L. pneumophila serogroup 1 strain, Paris strain and Lens strain found in different French départements (2) and low-level sequencing of covering of two strains not belonging to the L. pneumophila species. The strains selected are L. longbeachae, responsible for cases of legionellosis essentially in Australia, and L. anisa frequently found in water circulation but not found in patients.

The complete sequencing has likewise been carried out within the scope of the invention of an epidemic L. pneumophila serogroup 1 strain known as « Lens strain », responsible for a major epidemic in France with 86 cases and 17 deaths between November 2003 and January 2004.

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The genes found to be variable between the strains of Legionella, as well as preserved genes having a functional tie to the virulence of L.pneumophila can be utilized to manufacture DNA chips.

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A large number of isolates of various origins isolated from the environment or originating from clinical cases could be analyzed by using this tool to identify markers enabling the two categories of strains to be discriminated. The comparison of endemic and epidemic isolates will likewise provide bases for understanding the specificities of these strains and in particular the adaptability and stability of the Paris strain.

This approach also helps identify new functions necessary to the virulence of Legionella in humans and aid understanding of the different stages of this disease.

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The object of the invention is to allow development of novel tools for typing the strains of Legionella. These tools could be of the DNA "chip" type or of another type. The novel characteristics of these typing tools will be the following:

- * Rapidity and simplicity of use
- * High capacity for discrimination between the strains

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* Possibility for providing information on the genomic content of the strain analyzed and possibly prevent the risk associated with contamination by Legionella.

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The inventors have, during this study, brought to light genes found to be variable between the strains of Legionella, as well as preserved genes having a functional tie to wall or cellular envelope, or the virulence of L. pneumophila, genes which will be able to be used for carrying out these processes or diagnostic kits, especially for producing biochips with protein or DNA.

A large number of isolates of diverse origin isolated from the environment or originating from clinical cases could be analyzed by using these tools for the purpose of identifying markers for discriminating the two categories of strains.

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This approach will also help identify new functions necessary for the virulence of Legionella in humans and aid in comprehension of the different stages of this disease.

The object of the invention is thus to allow development of novel tools for typing of strains of Legionella. These tools could especially be of the protein or

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DNA/RNA biochip type. The novel characteristics of these typing tools will be the following:

- rapidity and simplicity of use;
- high capacity for discrimination between strains; and
- possibility of supplying information on the genomic content of the strain analyzed and of possibly preventing the risk associated with contamination by *Legionella*.

The inventors have, in the first instance, sequenced the complete genome of L. pneumophila Paris strain in the form of 56 contigs (SEQ ID No. 1 to 56), a sequence made up of a long chromosome of around 3.65 Mb and a long plasmid of around 36 kb. The inventors have likewise identified on these contigs (SEQ ID No. 1 to 56) the nucleic sequences coding for proteins with their respective function (cf Table I with the annotated sequences). The inventors have additionally compared these sequences to the sequences of the genome of L. pneumophila Philadelphia strain available at the website http://genome3.cpmc.columbia.edu/~legion/index.html and revealed their presence or not in the sequences of this Philadelphia strain. The sequences of the genome of L. pneumophila Philadelphia strain available on the website http://genome3.cpmc.columbia.edu/~legion/index.html correspond to the sequences of the 51 contigs identified in the list of sequences under the SEQ ID Nos. 3456 to 3506. This comparison, made from the available genomic sequence of L. pneumophila Philadelphia strain and the proteic sequences obtained by the inventors from the 6 possible reading frames, has revealed that some 88 % of these two genomes are very strongly preserved (95 to 100 % of proteic identity), the remaining 12 % being specific to each strain (cf. Tables I and IV). These results thus demonstrate that there is a wide genomic diversity within the L. pneumophila species. A serine protease autotransporter homolog in which is inserted ten repetitions in tandem of a pattern of 60 amino acids was especially identified among the genes specific to the Paris strain. The autotransporters are secretion systems for the Gram negative bacteria in which the Nterminal and C-terminal parts respectively enable secretion across the internal membrane and the formation of pores in the external membrane. The central part of the protein can then remain exposed at the level of the cellular surface or can be split and salted out in the external medium. The role of certain autotransporters in the virulence of the negative Gram bacteria has already been shown; furthermore, work on the autotransporters of the enterobacteria has helped identify a group of serine protease

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present solely in pathogenic bacterias whereof the diversity of functions could be linked to the adaptation to the niche occupied by the pathogen.

In addition, the inventors have revealed a very wide inter-species diversity from the sequencing of a pathogenic strain of *L. longbeachae*, which causes very few cases of legionellosis in France, but which is the major source of legionellosis in Australia, and that of a non-pathogenicic *L. anisa* strain. We were able to identify to date 703 ORFs of the *L. longbeachae* strain, whereof 53 % are specific to it; the majority of ORFs preserved between the *L. longbeachae* and *L. pneumophila* Paris strains have a percentage of proteic homology greater than 80%. The preliminary results obtained by the inventors on the non-pathogenic *L. anisa* strain have helped identify 54 % of specific sequences and un percentage of proteic homology greater than 70 % for the sequences preserved between the *L. anisa* and *L. pneumophila* Paris strains.

Tables I and II (« bestblast » obtained for each of the nucleic and proteic sequences corresponding to the annotated ORFs) and X to XXI hereinbelow comprise for each of the ORFs identified either in the Paris strain (Tables I and XIV), or in the Lens strain (Table XVI), its position on the contigs or chromosomes, and, if required, the existence of a peptide signal, the best result of the blast on nrprot (Best-Blastp). The ORFs:

- specific to the *L. pneumophila* Paris strain relative to the *L. pneumophila* Philadelphia strain;
- specific to the *L. pneumophila* Paris strain relative to the *L. pneumophila* Philadelphia and Lens strains (Table XVII);
- specific to the *L. pneumophila* Lens strain relative to the *L. pneumophila* Philadelphia and Paris strains (Table XVIII);
- specific to the *L. pneumophila* Philadelphia strain relative to the *L. pneumophila* Paris and Lens strains (Table XI);

were identified in considering as specific the ORFs having a percentage of proteic identity less than 75 %. In the event where ORF is preserved in the two genomes, the percentage of identity between the two proteins is mentioned.

In this Table I, the ORFs present in the partial sequence of the *L. longbeachae* strain have likewise been noted. Finally, the ORFs specific to the *Legionella* genre were identified by considering as specific the ORFs having a percentage of proteic identity with sequences of the nrprot bank less than 25 %.

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In Table XIX, the ORFs present at the same time in the Paris and Lens strain were indicated, though absent in the Philadelphia strain.

In Table XX, the ORFs present at the same time in the Paris and Philadelphia strain were indicated, though absent in the Lens strain.

Finally, in Table XXI, the ORFs present at the same time in the Philadelphia and Lens strain were indicated, though absent in the Paris strain.

In conclusion, the diversity revealed by the content of the present patent application helps define proteic probes, such as antibodies, or DNA probes for developing a typing tool. The utilization of this tool on a large number of strains isolated from patients and strains isolated from the environment will enable this tool to be validated, a tool which will aid in predicting the risk associated with a strain by discriminating in a certain manner the strains isolated from patients of other strains.

Among the significant families of proteins of Legionella pneumophila Paris strain the family of surface proteins or that of proteins implied in the biosynthesis of surface polysaccharides can be cited, or again that of proteins implied in the virulence of these bacteria. The process of evolution has allowed the development of a number of unique mechanisms on the Gram+bacteria, by which they can immobilize proteins on their surface. The functions of these different proteins of cellular walls are extremely diverse. However, many proteins linked covalently to the surface of the Gram+pathogens are estimated to be important for the survival of the pathogen inside the infected host. The study of Legionella pneumophila Paris strain demands novel approaches, in particular genetic, to improve understanding of the different metabolic paths of this organism.

Accordingly, it is object of the present invention to divulge the complete sequence of the genome of Legionella pneumophila Paris strain (Collection de the Institut Pasteur CIP 107-629-T), a sequence obtained from a collection of clones (BAC) filed on 19 November 2003 with the Collection Nationale de Cultures de Microorganisms (CNCM) [National Collection of Microorganism Cultures], 25 rue du Docteur Roux, 75724 Paris Cedex 15, France, according to the arrangements of the Budapest Treaty and registered under file number I-3137, as well as all the genes contained in said genome.

It is also another object of the present invention to divulge the complete sequence of the genome of *Legionella pneumophila* Lens strain, a sequence obtained from a collection of clones (BAC) filed on 23 September 2004 with the Collection

Nationale de Cultures de Microorganisms (CNCM) [National Collection of Microorganism Cultures], 25 rue du Docteur Roux, 75724 Paris Cedex 15, France, according to the arrangements of the Budapest Treaty and registered under file number I-3306, as well as all the genes contained in said genome.

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In effect, knowledge of the genome of these organisms enables the interactions between the different genes, the different proteins, and the different metabolic paths to be better defined. In effect, and contrary to divulging isolated sequences, the complete genomic sequence of an organism forms a whole, allowing all the information necessary to this organism to grow and function to be obtained immediately.

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If the present invention provides the nucleotidic sequence of the genome of Legionella pneumophila Paris strain (Collection de the Institut Pasteur CIP 107-629-T), having been the object of a filing of a collection of clones (BAC) covering this genome with the C.N.C.M. in Paris on 19 November 2003 and registered under file number I-3138, and likewise provides the nucleotidic sequence of the genome of Legionella pneumophila Lens strain, a sequence obtained from a collection of clones (BAC) filed on 23 September 2004 with CNCM, and likewise provides certain polypeptide sequences coded by these two genomes, the specialist will be able to determine the other ORFs, by utilizing known methods, and appropriate software.

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In the set of claims hereinbelow, the term «nucleotidic sequence» will especially be able to be replaced by the term «polynucleotide» without modifying the object and the scope of the set of claims such as filed.

The present invention thus relates to:

- a genomic nucleotidic sequence of *Legionella pneumophila* Paris strain characterized in that it is selected among the sequences SEQ ID 3507 and 3508, SEQ ID N° 55 and the sequences SEQ ID N° 1 to SEQ ID N° 54, and SEQ ID N° 56;
- a genomic nucleotidic sequence of *Legionella pneumophila* Lens strain characterized in that it is selected among the sequences SEQ ID 6733 and 6734.

The present invention likewise relates to an isolated or purified nucleotidic sequence:

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- (A) of Legionella pneumophila Paris strain, characterized in that it is selected among:
- a) a nucleotidic sequence comprising at least one sequence having 80 % identity with the sequences SEQ ID 3507 and 3508, and SEQ ID N $^\circ$ 1 to SEQ ID N $^\circ$ 56;

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- b) a nucleotidic sequence hybridizing in very stringent conditions with the sequences SEQ ID 3507 and 3508, and SEQ ID N° 1 to SEQ ID N° 56;
- c) a nucleotidic sequence complementing sequences SEQ ID 3507 and 3508 and SEQ ID N° 1 to SEQ ID N° 56, or complementing a nucleotidic sequence such as defined in a), or b), or a corresponding nucleotidic sequence of RNA; and
- d) a nucleotidic sequence of at least 15 nucleotides of fragment representative of sequences SEQ ID 3507 and 3508, and SEQ ID N° 1 to SEQ ID N° 56, or of fragment representative of their sequence, or
- (B) of Legionella pneumophila Lens strain, characterized in that it is selected among:
 - a) a nucleotidic sequence comprising at least one sequence having 80 % of identity with the sequences SEQ ID 6733 and 6734;
 - b) a nucleotidic sequence hybridizing in very stringent conditions with the sequences SEQ ID 6733 and 6734;
- c) a nucleotidic sequence complementing sequences SEQ ID 6733 and 6734; or complementing a nucleotidic sequence such as defined in a), or b), or a corresponding nucleotidic sequence of RNA; and
 - d) a nucleotidic sequence of at least 15 nucleotides of fragment representative of sequences SEQ ID 6733 and 6734; or of fragment representative of their sequence.

More particularly, the object of the present invention likewise is the nucleotidic sequences characterized in that they originate from sequences SEQ ID 3507 and 3508, and SEQ ID N° 1 to SEQ ID N° 56, and in that they code for a polypeptide selected from amongst the polypeptides of sequences SEQ ID 3509 to SEQ ID 6732, and SEQ ID N° 56 to SEQ ID N° 3455, preferably coding for a secreted enzyme likewise present in the Lens strain and Philadelphia strain of sequences SEQ ID 3675, 4267, 4292 and 6477, preferably coding for a polypeptide present on the surface of *Legionella pneumophila* Paris strain of sequence SEQ ID Nos. 3410, 704, 746, 2267, 2751, 3192, 3218, 3221, 3222, 3317, 3324, 136, 171, 310, 337, 481, 527 652, 664, 893, 972, 1148, 1298, 1361, 1503, 1521, 1576, 1651, 1755, 1847, 1877, 2224, 2406, 2843, 2930, 3037, 3139, 3157, 3165, 3181, preferably coding for a polypeptide present on the specific surface of *Legionella pneumophila* Paris strain relative to the Philadelphia strain, especially of sequence SEQ ID Nos. 3410, 171, 337, 481, 652, 1148, 1521, 2843, 3037, 3181, or one of its representative fragments of at least 5 amino acids, or coding for a

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polypeptide implied in the biosynthesis of polyssacharides having a cellular envelope of sequence SEQ ID Nos. 1126, 3218, 288, 632, 917, 1503, 1555, 1877, 1928, 1963, 2204, 2212, 2243, 2324, 2378, 2410, 2411, or again the nucleic sequences of the rtxA gene, especially those coding for the polypeptides of sequences SEQ ID Nos. 3410, 3037, 3165 and 3181.

The present invention also relates more generally to the nucleotidic sequences derived from the sequences SEQ ID Nos. 3507 and 3508, and SEQ ID N° 1 to SEQ ID N° 56, and coding for a polypeptide of *Legionella pneumophila* Paris strain such that they can be isolated from these sequences SEQ ID Nos. 3507 and 3508, and SEQ ID N° 1 to SEQ ID N° 56.

In addition, the nucleotidic sequences characterized in that they comprise a nucleotidic sequence selected among:

- a) a nucleotidic sequence coding for a polypeptide selected from amongst the sequences SEQ ID Nos. 3509 to 6732, and SEQ ID N° 56 to SEQ ID N° 3455;
- b) a nucleotidic sequence comprising at least 80 %, 85 %, 90 %, 95 % or 98 % of identity with a nucleotidic sequence coding for a polypeptide selected from amongst the sequences SEQ ID Nos. 3509 to 6732, and SEQ ID N° 56 to SEQ ID N° 3455;
 - c) a nucleotidic sequence hybridizing in very stringent conditions with a nucleotidic sequence coding for a polypeptide selected from amongst the sequences SEQ ID Nos. 3509 to 6732, and SEQ ID N° 56 to SEQ ID N° 3455;
 - d) a complementary nucleotidic sequence or RNA corresponding to a sequence such as defined in a), b) or c);
 - e) a nucleotidic sequence of a representative fragment having at least 15 nucleotides of a sequence such as defined in a) or d); and
- 25 f) a modified nucleotidic sequence of a sequence such as defined in a), d) or e), are likewise objects of the invention.

Nucleic acid, nucleic sequence or nucleic acid, polynucleotide, oligonucleotide, polynucleotidic sequence, nucleotidic sequence, terms which will be employed indifferently in the present description, are understood to designate precise chaining of nucleotides, modified or not, effectively defining a fragment or a region of a nucleic acid, comprising or not non-natural nucleotides, and able to correspond just as well to a double-strand DNA, a single-strand DNA as transcription products of said DNAs. Therefore, the nucleic sequences according to the invention likewise encompass the PNA (Peptid Nucleic Acid), or similar.

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It must be understood that the present invention does not relate to the nucleotidic sequences in their natural chromosomic environment, that is, in the natural state. These are sequences which were isolated and/or purified, that is they were sampled directly or indirectly, for example by copy, their environment having been at least partially modified. It is understood to likewise designate the nucleic acids obtained by chemical synthesis.

« Percentage of identity » between two sequences of nucleic acids or amino acids in the sense of the present invention is understood to designate a percentage of nucleotides or residues of identical amino acids between the two sequences to be compared, obtained after the best alignment, this percentage being purely statistical and the differences between the two sequences being distributed randomly and over their entire length. "Best alignment" or " optimal alignment " is understood to designate the alignment for which the percentage of identity determined hereinafter is the highest. The comparisons of sequences between two sequences of nucleic acids or amino acids are traditionally made by comparing these sequences after they were aligned in optimal fashion, said comparison being made by segment or by « window of comparison » to identify and compare the local regions of similarity of sequence. The optimal alignment of the sequences for comparison can be made, apart from manually, by means of the local of de Smith and Waterman (1981, Ad. App. Math. 2:482), by means of the local homology algorithm of Neddleman and Wunsch (1970, J. Mol. Biol. 48:443), by means of the similarity search method of Pearson and Lipman (1988, Proc. Natl. Acad. Sci. USA 85:2444), by means of software utilizing these algorithms (GAP, BESTFIT, BLAST P, BLAST N, FASTA and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer Group, 575 Science Dr., Madison, WI). To obtain optimal alignment, the program BLAST is preferably used, with the BLOSUM 62 matrix. The PAM or PAM250 matrices can likewise be used.

The percentage of identity between two sequences of nucleic acids or amino acids is determined by comparing these two sequences aligned optimally in which the sequence of nucleic acids or amino acids to be compared can comprise additions or deletions relative to the reference sequence for optimal alignment between these two sequences. The percentage of identity is calculated by determining the number of identical positions for which the nucleotide or the residue of amino acid is identical between the two sequences, by dividing this number of identical positions by the total

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number of positions compared and by multiplying the result obtained by 100 to obtain the percentage of identity between these two sequences.

Nucleic sequences having a percentage of identity of at least 80 %, preferably 85 % or 90 %, more preferably 95 % or 98 % or 99 %, after optimal alignment with a reference sequence are understood to designate he nucleic sequences exhibiting, relative to the nucleic reference sequence, certain modifications such as in particular deletion, truncation, elongation, chimeric fusion and/or substitution, especially specific, and whereof the nucleic sequence has at least 80 %, preferably 85 %, 90 %, 95 %, 98 % or 99 %, of identity after optimal alignment with the nucleic reference sequence. These are preferably sequences whereof the complementary sequences are capable of being hybridized specifically with the reference sequences. Preferably, the specific hybridization conditions or stringent conditions will be such that they ensure at least 80 %, preferably 85 %, 90 %, 95 %, 98 % or 99 % of identity after optimal alignment between one of the two sequences and the sequence complementary to the other.

Hybridization in very stringent conditions signifies that the temperature and ionic force conditions are selected in such a way that they permit hybridization to be maintained between two fragments of complementary DNA. By way of illustration, very stringent conditions of the hybridization stage for the purpose of defining the polynucleotide fragments described hereinabove, are advantageously the following.

DNA-DNA or DNA-RNA hybridization is performed in two stages: (1) prehybridization at 42°C for 3 hours in phosphate buffer (20 mM, pH 7.5) containing 5 x SSC (1 x SSC corresponds to a solution 0.15 M NaCl + 0.015 M sodium citrate), 50 % formamide, 7 % sodium dodecyl sulfate (SDS), 10 x Denhardt's, 5 % dextran sulfate and 1 % DNA salmon sperm; (2) hybridization per se for 20 hours at a temperature depending on the size of the probe (i.e.: 42°C, for a probe of size > 100 nucleotides) followed by 2 washes of 20 minutes at 20°C in 2 x SSC + 2 % SDS, 1 wash of 20 minutes at 20°C in 0.1 x SSC + 0.1 % SDS. The last wash is done in 0.1 x SSC + 0.1 % SDS for 30 minutes at 60°C for a probe of size > 100 nucleotides. The very stringent hybridization conditions described hereinabove for a polynucleotide of defined size can be adapted by the specialist for oligonucleotides of larger or smaller size, according to the teaching of Sambrook *et al.*, (1989, Molecular cloning: a laboratory manual. 2nd Ed. Cold Spring Harbor).

In addition, fragment representative of sequences according to the invention is understood to designate any nucleotide fragment having at least 15 nucleotides,

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preferably at least 20, 25, 30, 50, 75, 100, 150, 300 and 450 consecutive nucleotides of the sequence from which it originates.

Representative fragment is understood in particular to be a nucleic sequence coding for a biologically active fragment of a polypeptide, such as defined hereinbelow.

Representative fragment is likewise understood to be the intergenic sequences, and in particular the nucleotidic sequences bearing the regulation signals (promoters, terminators, or enhancers, ...), or again probe or primer sequences aiding in specifically detecting or amplifying the nucleic sequences coding for the polypeptides of sequences SEQ ID Nos. 3509 to 6732, and SEQ ID N° 56 to SEQ ID N° 3455.

Of said representative fragments those are preferred having nucleotidic sequences corresponding to open reading frames, known as ORF sequences (ORF for « Open Reading Frame »), included in general between an initiation codon and a stop codon, or between two stop codons, and coding for polypeptides, preferably of at least 100 amino acids, such as for example, without limiting them, the ORF sequences to be described hereinafter.

The representative fragments according to the invention can be obtained for example by specific amplification such as PCR or after digestion by appropriate restriction enzymes of nucleotidic sequences according to the invention, this method being described in particular in the work of Sambrook *et al.*. Said representative fragments can likewise be obtained by chemical synthesis as long as their size is not too significant, according to methods well known to the specialist.

The representative genome fragments of Legionella pneumophila Paris strain according to the invention likewise comprise at least one fragment of at least 15 nucleotides or more as cited above for the fragments resulting from enzymatic cutting at the level of a restriction site. Of course, expression proteins such as RNA or proteins are understood according to the present invention.

Among the sequences containing inventive sequences, or representative fragments, are likewise understood the sequences which are naturally framed by sequences which present at least 80 %, 85 %, 90 %, 95 % or 98 % of identity with the sequences according to the invention.

Modified nucleotidic sequence is understood as any nucleotidic sequence obtained by mutagenesis according to techniques well known to the specialist, and comprising des, preferably a maximum 10 %, 7.5 %, 5 %, 2.5 %, 1 %, 0.5 %, 0.1 % or even less than 0.01 %, of modified nucleotides, relative to normal sequences, for

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example mutations in the regulating and/or promoting sequences of the expression of the polypeptide, especially leading to modification of the rate of expression or activity of said polypeptide.

Modified nucleotidic sequence is likewise understood as any nucleotidic sequence coding for a polypeptide modified such as defined hereinbelow.

The representative fragments according to the invention can likewise be probes or primers, which can be utilized in processes for detection, identification, dosage or amplification of nucleic sequences.

In a preferred manner the invention is relative to a nucleotidic sequence coding for a polypeptide according to the invention.

In a preferred manner the invention is relative to a nucleotidic sequence according to the invention, characterized in that it codes for a specific polypeptide of a bacteria of the *Legionella* genre, or one of its fragments of at least 5 amino acids, or its complementary nucleic sequence.

In a preferred manner the invention is relative to a nucleotidic sequence according to the invention, characterized in that it codes for a specific polypeptide of a pathogenic bacteria of the *Legionella* genre and/or of the *Legionella* pneumophila species, or one of its fragments of at least 5 amino acids, or its complementary nucleic sequence.

In a preferred manner the invention is relative to a nucleotidic sequence according to the invention, characterized in that it codes for a specific polypeptide of a bacteria of the *Legionella pneumophila* species Paris strain, or one of its fragments of at least 5 amino acids, or its complementary nucleic sequence.

In a preferred manner the invention is relative to a nucleotidic sequence according to the invention, characterized in that it codes for a specific polypeptide of a bacteria of the *Legionella pneumophila* species Paris strain relative to the Philadelphia strain, or one of its fragments of at least 5 amino acids, in particular selected from amongst the polypeptides of sequence SEQ ID Nos. 3410, 171, 337, 481, 652, 1148, 1521, 2843, 3037, 3181 or one of its fragments of at least 5 amino acids, or its complementary nucleic sequence.

In a preferred manner the invention is relative to a nucleotidic sequence according to the invention, characterized in that it codes for a specific polypeptide of a bacteria of the *Legionella pneumophila* species Paris strain relative to the Lens and Philadelphia strains, or one of its fragments of at least 5 amino acids, in particular

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selected from amongst the polypeptides whereof the sequences are indicated in Table XVII or one of their fragments of at least 5 amino acids, or their complementary nucleic sequence.

In a preferred manner the invention is relative to a nucleotidic sequence according to the invention, characterized in that it codes for a surface polypeptide of *Legionella pneumophila* Paris strain, or one of its fragments of at least 5 amino acids, in particular selected from amongst the sequence polypeptides SEQ ID Nos. 3410, 704, 746, 2267, 2751, 3192, 3218, 3221, 3222, 3317, 3324, 136, 171, 310, 337, 481, 527 652, 664, 893, 972, 1148, 1298, 1361, 1503, 1521, 1576, 1651, 1755, 1847, 1877, 2224, 2406, 2843, 2930, 3037, 3139, 3157, 3165, 3181, or one of its fragments of at least 5 amino acids, or its complementary nucleic sequence.

In a preferred manner the invention is relative to a nucleotidic sequence according to the invention, characterized in that it codes for a polypeptide of specific surface of *Legionella pneumophila* Paris strain relative to the Philadelphia strain, selected from amongst the polypeptides of sequences SEQ ID Nos. 3410, 171, 337, 481, 652, 1148, 1521, 2843, 3037, 3181, or one of its fragments of at least 5 amino acids, or its complementary nucleic sequence.

In a preferred manner the invention is relative to a nucleotidic sequence according to the invention, characterized in that it codes for a polypeptide implied in the biosynthesis of polysaccharide having a cellular envelope of *Legionella pneumophila* Paris strain, in particular selected from amongst the polypeptides of sequence SEQ ID Nos. 1126, 3218, 288, 632, 917, 1503, 1555, 1877, 1928, 1963, 2204, 2212, 2243, 2324, 2378, 2410, 2411, or one of its representative fragments of at least 5 amino acids, or its complementary nucleic sequence.

In a preferred manner the invention is relative to a nucleotidic sequence according to the invention, characterized in that it codes for a polypeptide of *Legionella pneumophila* Paris strain coded by the rtxA gene of sequence SEQ ID Nos. 3410, 3037, 3165 and 3181, or one of its representative fragments of at least 5 amino acids, or its complementary nucleic sequence.

The rtxA gene of Legionella pneumophila was demonstrated as being implied in virulence. This gene codes for a protein of 1208 aa. Four ORFs whereof the references are given hereinbelow (SEQ ID 3410 - 3037 - 3165 - 3181) correspond to this gene in the Paris strain which would code for a protein of at least 4000 aa. Comparison with the Philadelphia strain shows the presence of a homologous gene for the N- and C-terminal

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parts. The central part of the protein is constituted by repetitions of a pattern of around 20 aa different between the two strains.

In another aspect, in a preferred manner, the invention is relative to a nucleotidic sequence according to the invention characterized in that it codes for a polypeptide of *Legionella pneumophila* Paris strain, or one of its representative fragments of at least 5 amino acids implied in the biosynthesis of the amino acids, in the biosynthesis of the cofactors, prosthetic and transporter groups, implied in the cellular machinery, implied in the central intermediate metabolism, implied in the energetic metabolism, implied in the metabolism of fatty acids and phospholipids, implied in the metabolism of nucleotides, purines, pyrimidines or nucleosides, implied in the functions of regulation, implied in the process of transcription, implied in the process of transcription, implied in the process of translation, implied in the process of translation to atypical conditions, implied in sensitivity to medications and the like, or implied in the functions relative to transposons.

Owing to the genomic sequence presented in the present invention, the specialist will know how to identify the genes coding for proteins regulating transcription of the genes of *Legionella pneumophila* Paris strain. In addition, Table I provides the list of the open reading phases (ORF for «Open Reading Frame) annotated and identified on the genome of *Legionella pneumophila* Paris strain (SEQ ID N° 1 to SEQ ID N° 55), with especially their position on said genome, and, in Table II, the putative functions which can be attributed to them by utilizing customary techniques for comparing the genomic («Bestblast »). All the same, such a list must not be considered as limiting, where one protein can have several roles in the cell.

Modifying the structure or the integrity of these genes could help modify the expression of the target genes controlled by the target promoters of these regulators. Thus, the expert will be able to select the regulator(s) pertinent for the required application as well as their target, thus allowing optimization of the expression of genes of interest. The utilization of the tools described above such as the DNA chips, also registers all the genes whereof the regulation is modified by inactivation of certain genes. It is thus possible to select a set of control sequences responding to the same type of regulation. These sequences can then be used to control the expression of genes of interest.

In general, the list of sequences SEQ ID, or their corresponding coding nucleic sequence could be determined by the specialist from the most probable putative

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functions determined for each of the sequences SEQ ID in Tables I and XIV hereinbelow for each of the classes of activity classified hereinbelow.

It is important to note all the same that a living organism is a whole and must be taken as such. Accordingly, so as to develop and exhibit its properties, any organism has need of interaction between the different metabolic paths. Therefore, the abovementioned classification must not be considered as limiting, with a gene able to be implied in two distinct metabolic paths.

The invention likewise relates to polypeptides coded by a nucleotidic sequence according to the invention, preferably by a fragment representative of the sequences SEQ ID Nos. 3509 to 6732, the sequence SEQ ID N° 55 or sequences SEQ ID N° 1 to SEQ ID N° 54, and SEQ ID N° 56, and corresponding to an ORF sequence, such as described in Table XIV (coding for one of the sequences SEQ ID N° 3509 to SEQ ID N° 6732), and in Table I (coding for one of the sequences SEQ ID N° 57 to SEQ ID N° 3455).

- In particular, polypeptides of *Legionella pneumophila* Paris strain, characterized in that they are selected from amongst the following polypeptides:
 - polypeptides of sequences SEQ ID Nos. 3509 to 6732 and of sequences SEQ ID N° 56 to SEQ ID N° 3455;
 - preferably enzymes secreted by *Legionella pneumophila* Paris, Lens and Philadelphia strains, especially sequences SEQ ID Nos. 3675, 4267, 4292 and 6477;
 - preferably polypeptides present on the surface of *Legionella pneumophila* Paris strain, especially of sequence SEQ ID Nos. 3410, 704, 746, 2267, 2751, 3192, 3218, 3221, 3222, 3317, 3324, 136, 171, 310, 337, 481, 527 652, 664, 893, 972, 1148, 1298, 1361, 1503, 1521, 1576, 1651, 1755, 1847, 1877, 2224, 2406, 2843, 2930, 3037, 3139,
- 3157, 3165, 3181, and still more preferred polypeptides present on the specific surface of Legionella pneumophila Paris strain relative to the Philadelphia strain, especially those of sequences SEQ ID Nos. 3410, 171, 337, 481, 652, 1148, 1521, 2843, 3037, 3181, or one of its representative fragments of at least 5 amino acids;
- polypeptides implied in the biosynthesis of polysaccharides having a cellular envelope, especially of sequence SEQ ID Nos. 1126, 3218, 288, 632, 917, 1503, 1555, 1877, 1928, 1963, 2204, 2212, 2243, 2324, 2378, 2410, 2411;
 - or again polypeptides of sequence SEQ ID Nos. 3410, 3037, 3165 and 3181, coded by the rtxA gene.

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The invention likewise comprises polypeptides characterized in that they comprise a polypeptide selected from amongst:

- a) a polypeptide of sequences SEQ ID Nos. 3509 to 6732, and of sequences SEQ ID N° 56 to SEQ ID N° 3455;
- b) a polypeptide having at least 80 % preferably 85 %, 90 %, 95 % and 98 % of identity with an polypeptide of sequences SEQ ID Nos. 3509 to 6732, and of sequences SEQ ID N° 56 to SEQ ID N° 3455;
 - c) a fragment of at least 5 amino acids, preferably biologically active, of one such as defined in b);
- d) a biologically active fragment of a polypeptide of sequence SEQ ID N° 56 to SEQ ID N° 3455; and
 - e) a modified polypeptide of a polypeptide of sequences SEQ ID Nos. 3509 to 6732, and of sequences SEQ ID N° 56 to SEQ ID N° 3455, comprising at most 10 % modified amino acids, preferably 7.5 %, 5 %, 2.5 %, 1 %, 0.5 %, 0, 1 % or again 0.01 %.

The nucleotidic sequences coding for the abovedescribed polypeptides are likewise an object of the invention.

In the present description, the terms polypeptides, polypeptide sequences, peptides and proteins are interchangeable.

It must be understood that the invention does not relate to polypeptides in the natural form, that is, that they are not taken in their natural environment, rather they were isolated or obtained by purification from natural sources, or else obtained by genetic recombination, or by chemical synthesis, and that they can then comprise non-natural amino acids such as will be described hereinbelow.

Polypeptide having a certain percentage of identity with another, likewise designated by homologous polypeptide, is understood to designate those polypeptides having, relative to natural polypeptides, certain modifications, in particular deletion, addition or substitution of at least one amino acid, truncation, elongation, a chimeric solution and/or a mutation, or polypeptides having post-translational modifications. Of the homologous polypeptides preference is given to those whereof the sequence of amino acids has at least 80 %, preferably 85 %, 90 %, 95 %, 98 % and 99 % of identity with the sequences of amino acids of the polypeptides according to the invention. In the case of substitution, one or more consecutive or non-consecutive amino acid(s) are replaced by « equivalent » amino acids. The expression « equivalent amino acids» in

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this case endeavors to designate any amino acid likely to be substituted for one of the amino acids of the base structure without however essentially modifying the biological activities of the corresponding peptides and such as they will be defined hereinbelow.

These equivalent amino acids can be determined either by being supported on their structural homology with the amino acids for which they are substituted, or on results of comparative assays of biological activity between the different polypeptides capable of being effected.

By way of example, mention is made of the possibilities of substitution for being carried out without the resulting extensive modification of the biological activity of the corresponding modified polypeptide. Thus leucine can be replaced by valine or isoleucine, aspartic acid by glutamine acid, glutamine by asparagine, arginine by lysine, etc., the inverse substitutions naturally being envisaged under the same conditions.

The homologous polypeptides correspond likewise to the polypeptides coded by the homologous or identical nucleotidic sequences, such as defined previously and thus comprise in the present definition mute polypeptides or corresponding to inter or intra species variations, able to exist in *Legionella*, and which correspond especially to truncations, substitutions, deletions and/or additions, of at least one residue of amino acids.

It is understood that the percentage of identity between two polypeptides is calculated in the same way as between two sequences of nucleic acids. Thus, the percentage of identity between two polypeptides is calculated after optimal alignment of these two sequences, on a maximum homology window. To define said maximum homology window, the same algorithms as for the nucleic acid sequences can be utilized.

Biologically active fragment of a polypeptide according to the invention is understood to mean in particular a fragment of polypeptide, such as defined hereinbelow, having at least one of the biological characteristics of the polypeptides according to the invention, especially in that it capable of exerting in general an even partial activity, such as for example:

- enzymatic (metabolic) activity or an activity able to be implied in the biosynthesis or biodegradation of organic or inorganic compounds;
 - structural activity (cellular envelope, coping molecule, ribosome);
 - transport activity (energy, ion); or in the secretion of protein;

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- activity in the process of replication, amplification, preparation, transcription, translation or maturation, especially of DNA, RNA or proteins.

Fragment of polypeptides according to the invention is understood to mean a polypeptide comprising a minimum of 5 amino acids, preferably 6, 7, 8, 9, 10, 15, 20, 25, 30, 40, 50, 75, 100, 150 acids and 200 amino acids.

The fragments of polypeptides can correspond to isolated or purified fragments naturally present in the strains of *Legionella*, or to fragments which can be obtained by cleavage of said polypeptide by a proteolitic enzyme such as trypsine or chymotrypsine or collagenase, by a chemical reagent (cyanogen bromide, CNBr) or by placing said polypeptide in a highly acidic environment (for example at pH = 2.5). Polypeptidic fragments can likewise be prepared by chemical synthesis, from hosts transformed by a vector of expression according to the invention which contains a nucleic acid allowing expression of said fragment, and placed under the control of the appropriate elements of regulation and/or expression.

« Modified polypeptide» of a polypeptide according to the invention is understood to mean a polypeptide obtained by genetic recombination or by chemical synthesis such as described below, which has at least one modification relative to the normal sequence, preferably at most 10 % of amino acids modified relative to the normal sequence, preferably even at most 7.5 %, 5 %, 2.5 %, 1 %, 0.5 %, 0, 1 % or again 0.01 %. These modifications can be especially made to amino acids necessary for the specificity or efficacy of the activity, or at the origin of the structural conformation, the charge, or the hydrophobicity of the polypeptide according to the invention. Polypeptides of equivalent, augmented or diminished activity, or of equivalent, narrower or wider specificity can thus be created. Among the polypeptides modified, those polypeptides in which up to five amino acids can be modified, truncated at the N end or C terminal, or else deleted, or added, must be mentioned.

As is indicated, the object of the modifications of a polypeptide especially are:

- to permit its usage in biosynthesis or biodegradation processes of organic or inorganic compounds,
- to permit its usage in processes of replication, amplification, repair and transcription, translation, or maturation especially of DNA, RNA, or proteins,
 - to permit its improved secretion,
 - to modify its solubility, efficacy or specificity of activity, or again to facilitate its purification.

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Chemical synthesis likewise has the advantage of being able to utilize nonnatural amino acids or non-peptidic bonds. Therefore, it can be interesting to utilize non-natural amino acids, for example in the D form, or analogs of amino acids, especially sulphurized forms.

In another aspect, the invention is preferably relative to a polypeptide according to the invention, characterized in that it is a specific polypeptide of a bacteria of the *Legionella* genre, or one of its fragments of at least 5 amino acids.

In another aspect, the invention is preferably relative to a polypeptide according to the invention, characterized in that it is a specific polypeptide of a pathogenic bacteria of the *Legionella* genre and/or of the *Legionella* pneumophila species, or one of its representative fragments of at least 5 amino acids.

In another aspect, preferably, the invention is relative to a polypeptide according to the present invention, characterized in that it is a specific polypeptide of a bacteria of the species *Legionella pneumophila* Paris strain, or one of its representative fragments of at least 5 amino acids.

In another aspect, preferably, the invention is relative to a polypeptide according to the present invention, characterized in that it is a specific polypeptide of a bacteria of the *Legionella pneumophila* species Paris strain relative to the Philadelphia strain, or one of its representative fragments of at least 5 amino acids, in particular selected from amongst the polypeptides of sequences SEQ ID Nos. 3410, 171, 337, 481, 652, 1148, 1521, 2843, 3037, 3181, or one of its representative fragments of at least 5 amino acids.

In another aspect, preferably, the invention is relative to a polypeptide according to the present invention, characterized in that it is a specific polypeptide of a bacteria of the *Legionella pneumophila* species Paris strain relative to the Lens and Philadelphia strains, or one of its representative fragments of at least 5 amino acids, in particular selected from amongst the polypeptides whereof the sequence is indicated in Table XVII, or one of their representative fragments of at least 5 amino acids.

In another aspect, preferably, the invention is relative to a polypeptide according to the present invention, characterized in that it is a surface polypeptide of *Legionella pneumophila* Paris strain, or one of its representative fragments of at least 5 amino acids, in particular selected from amongst the polypeptides of sequence SEQ ID Nos. 3410, 704, 746, 2267, 2751, 3192, 3218, 3221, 3222, 3317, 3324, 136, 171, 310, 337, 481, 527 652, 664, 893, 972, 1148, 1298, 1361, 1503, 1521, 1576, 1651, 1755, 1847,

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1877, 2224, 2406, 2843, 2930, 3037, 3139, 3157, 3165, 3181, or one of its representative fragments of at least 5 amino acids.

In another aspect, preferably, the invention is relative to a polypeptide according to the present invention, characterized in that it is a polypeptide of specific surface of *Legionella pneumophila* Paris strain relative to the Philadelphia strain, selected from amongst the polypeptides of sequence SEQ ID Nos. 3410, 171, 337, 481, 652, 1148, 1521, 2843, 3037, 3181, or one of its representative fragments of at least 5 amino acids.

In another aspect, preferably, the invention is relative to a polypeptide according to the invention, characterized in that it is a polypeptide *Legionella pneumophila* Paris strain implied in the biosynthesis of polysaccharide having a cellular envelope of *Legionella pneumophila* Paris strain, in particular selected from amongst the polypeptides of sequence SEQ ID Nos. 1126, 3218, 288, 632, 917, 1503, 1555, 1877, 1928, 1963, 2204, 2212, 2243, 2324, 2378, 2410, 2411, or one of its representative fragments of at least 5 amino acids.

In another aspect, preferably, the invention is relative to a polypeptide according to the invention, characterized in that it is a polypeptide de *Legionella pneumophila* Paris strain coded by the rtxA gene de sequence SEQ ID Nos. 3410, 3037, 3165 and 3181, or one of its representative fragments of at least 5 amino acids, or its complementary nucleic sequence.

In another aspect, preferably, the invention is relative to a polypeptide according to the invention, characterized in that it is a polypeptide of *Legionella pneumophila* Paris strain, or one of its representative fragments of at least 5 amino acids, implied in the biosynthesis of amino acids, in the biosynthesis of cofactors, prosthetic and transporter groups, implied in the cellular machinery, implied in the central intermediary metabolism, implied in the energetic metabolism, implied in the metabolism of fatty acids and phospholipids, implied in the metabolism of nucleotides, purins, pyrimidins or nucleosides, implied in the functions of regulation, implied in the process of replication, implied in the process of transcription, implied in the process of translation, implied in the process of transport and binding of proteins, implied in the adaptation to atypical conditions, implied in the sensitivity to drugs and the like, or implied in the functions relatives to transposons.

The object of the present invention is likewise the nucleotidic sequences and/or polypeptides according to the invention, characterized in that said sequences are registered on a registration support whose form and nature facilitate reading, analysis

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and/or exploitation of said sequence(s). These supports can likewise contain other information extracted from the present invention, especially the analogies with already known sequences, and/or information concerning the nucleotidic sequences and/or polypeptides of other microorganisms for the purpose of facilitating comparative analysis and exploitation of the results obtained.

Among said registration supports, preference is given in particular to those supports readable by a computer, such as magnetic, optic, electric or hybrid supports, in particular information discs, CD-ROM, servers. Such registration supports are likewise an object of the invention.

The registration supports according to the invention, with the information contributed, are very useful for the choice of primers or nucleotidic probes for determining genes in *Legionella pneumophila* Paris strain or strains close to this organism. Similarly, the utilization of these supports for the study of genetic polymorphism of a strain close to *Legionella pneumophila* Paris strain, in particular by determination of the colinearity regions, is very useful as far as these supports provide not only the nucleotidic sequence of the genome of *Legionella pneumophila* Paris strain, but likewise the genomic organization in said sequence. Thus, the uses of registration supports according to the invention are likewise objects of the invention.

The homology analysis between different sequences is completed in effect advantageously by means of software for sequence comparisons, such as BlastP or BlastN software, or other software well known to the specialist.

A probe or primer is defined, in the sense of the invention, as being a fragment of single-strand nucleic acids or a denatured double-strand fragment comprising for example 12 bases at several kb, especially 15 at several hundreds of bases, preferably from 15 to 50 or 100 bases, and possess a hybridization specificity in conditions determined to form a hybridization complex with a nucleic acid target.

The probes and primers according to the invention can be marked directly or indirectly by a radioactive or non-radioactive compound using methods well known to the specialist, to obtain a detectable and/or quantifiable signal.

The non-marked sequences of polynucleotides according to the invention can be utilized directly as probe or primer.

The sequences are generally marked to obtain sequences utilizable for numerous applications. The marking of the primers or probes according to the invention is done by radioactive elements or by non-radioactive molecules.

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Examples of the radioactive isotopes used are ³²P, ³³P, ³⁵S, ³H or ¹²⁵I. The non-radioactive entities are selected among ligands such as biotin, avidin, streptavidin, dioxygenin, haptenes, colorants, luminescent agents such as radioluminescent, chemoluminescent, bioluminescent, fluorescent, phosphorescent agents.

The polynucleotides according to the invention can thus be utilized as primer and/or probe in processes in particular making use of the PCR technique (amplification in chain by polymerase) (Rolfs *et al.*, 1991, Berlin: Springer-Verlag). This technique requires the choice of pairs of oligonucleotide primers framing the fragment to be amplified. For example, reference can be made to the technique described in the U.S. patent N° 4,683,202. The amplified fragments can be identified, for example after electrophoresis in agarose or polyacrylamide gel, or according to a chromatographic technique such as filtration on gel or ion exchange chromatography, then sequenced. The specificity of the amplification can be controlled by using as primer the inventive nucleotidic sequences of polynucleotides as matrix, plasmids containing these sequences or even the derived amplification products. The amplified nucleotide fragments can be utilized as reagents in hybridization reactions so as to reveal the presence, in a biological sample, of a nucleic acid target of sequence complementary to those of said amplified nucleotide fragments.

The aim of the invention is likewise the nucleic acids capable of being obtained by amplification by means of primers according to the invention.

Other amplification techniques for the nucleic acid target can be advantageously employed as an alternative to PCR (PCR-like) by means of a couple of primers of nucleotidic sequences according to the invention. PCR-like is understood to mean all the methods making use of direct or indirect reproductions of the sequences of nucleic acids, or else in which the marking systems were amplified; these techniques are well known, and in general are amplification of DNA by a polymerase; when the original sample is a RNA reverse transcription should be previously carried out. There are currently numerous processes enabling this amplification, such as for example the SDA technique (Strand Displacement Amplification) or brine displacement amplification technique (Walker *et al.*, 1992, Nucleic Acids Res. 20:1691), the TAS technique (Transcription-based Amplification System) described by Kwoh *et al.* (1989, Proc. Natl. Acad. Sci. USA, 86:1173), the 3SR technique (Self-Sustained Sequence Replication) described by Guatelli *et al.* (1990, Proc. Natl. Acad. Sci. USA 87:1874), the NASBA technique (Nucleic Acid Sequence Based Amplification) described by Kievitis *et al.*

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(1991, J. Virol. Methods, 35:273), the TMA technique (Transcription Mediated Amplification), the LCR technique (Ligase Chain Reaction) described by Landegren *et al.* (1988, Science 241:1077), the RCR technique (Repair Chain Reaction) described by Segev (1992, Kessler C. Springer Verlag, Berlin, New-York, 197-205), the CPR technique (Cycling Probe Reaction) described by Duck *et al.* (1990, Biotechniques, 9:142), the Q-beta-replicase amplification technique described by Miele *et al.* (1983, J. Mol. Biol., 171:281). Certain of these techniques have since been refined.

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In the event where the polynucleotide target to be detected is a RNAm, prior to use of an amplification reaction by means of the primers according to the invention or prior to use of a detection process by means of the inventive probes, an enzyme of inverse transcriptase type is used advantageously in order to obtain a DNAc from the RNAm contained in the biological sample. The DNAc obtained will then serve as target for the primers or the probes used in the process of amplification or detection according to the invention.

The technique of hybridization of probes can be executed in various ways (Matthews et al., 1988, Anal. Biochem., 169:1-25). The most general method consists of immobilizing the nucleic acid extracted from cells of different tissues or cells in culture on a support (such as nitrocellulose, nylon, polystyrene) and of incubating, in well-defined conditions, the nucleic acid target immobilized with the probe. After hybridization, the probe excess is eliminated and the hybrid molecules formed are detected by the appropriate method (measuring of radioactivity, fluorescence or enzymatic activity associated with the probe).

In accordance with another operating mode of the nucleic probes according to the invention, the latter can be utilized as capture probes. In this case, a probe, known as «capture probe», is immobilized on a support and serves to capture via specific hybridization the nucleic acid target obtained from the biological sample to be tested and the nucleic acid target is then detected due to a second probe, known as «detection probe», marked by an easily detectable element.

Of the possibly interesting fragments of nucleic acids, anti-sense oligonucleotides should thus be cited in particular, that is, whereof the structure ensures, via hybridization with the sequence target, inhibition of the expression of the corresponding product. Sense oligonucleotides which, by interaction with proteins implied in regulating the expression of the corresponding product, will cause either inhibition or activation of this expression, should likewise be cited.

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In a preferred way the probes or primers according to the invention are immobilized on a support, covalently or non-covalently. In particular, the support can be a DNA chip or a high-density filter, likewise objects of the present invention.

The interest in using DNA chips or, if required, protein chips in the domain of diagnostics or epidemiology rests as mentioned previously on the possibility of analyzing a large number of sequences at the same time and very rapidly, for:

- classification or typing of bacteria as a function of the presence of a sequence or of a profile of sequences characteristic of the genre, especially of the pathogenicity or not of the genre, especially Legionella, or of the species, especially Legionella pneumophila, or specific to a bacteria of the Legionella genre and/or of the Legionella pneumophila subspecies Paris strain, or specific to a bacteria of the Legionella pneumophila subspecies Paris strain relative to the Philadelphia strain and/or Lens strain, or even specific to a bacteria of the Legionella pneumophila subspecies Paris and Lens strain relative to the Philadelphia strain, in particular, in association with the gravity or not of the pathologies which such bacteria can induce in case of infection in mammals, especially in humans; or

- simultaneous comparison of sequence or of profile of sequences between different genres, species or strain of bacteria, pathogenic or not, allowing especially identification of a gene, or the corresponding proteic sequence, or a profile of genes whereof the presence and/or the expression in a bacteria is specific according to its genre, its species or its sub-species or strain of bacteria, and/or its pathogenicity or not. This information is largely useful especially for rapidly identifying the presence or not of a pathogenic bacteria, the gravity of the infection it can cause, the treatment adapted to an infection, and/or the necessity and the means for implementing contaminated circuits or fluids or able to be contaminated for decontaminating the objects. This information will likewise be largely useful to epidemiological studies relative to this genre of bacteria.

DNA chip or high-density filter is understood to mean a support on which DNA sequences are fixed, each of them able to be marked by its geographic location. These chips or filters differ principally in their size, the material of the support, and possibly the number of DNA sequences fixed thereto.

The probes or primers according to the first invention can be fixed on solid supports, in particular the DNA chips, by means of different fabrication processes. In

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particular, *in situ* synthesis can be carried out by photochemical addressing or via ink jet. Other techniques consist of carrying out *ex situ* synthesis and fixing the probes on the support of the DNA chip by mechanical or electronic addressing, or by ink jet. These different processes are well known to the specialist.

In effect, numerous techniques or devices for analysis of biological samples were developed in recent years, in particular for parallel analysis of several quantities of nucleic acids, especially following the development of the genomic.

Among these techniques or devices, the supports enabling high-rate analysis of nucleic acids, such as biochips, or DNA chips (also called « micro- or macroarrays », or even « DNA chip ») were the object of numerous studies.

These biochips can be made in particular from a support, generally solid and functionalized, on which given nucleic acids (nucleic probes) were fixed by covalent bond and localized, and on which nucleic probes are fixed specifically respectively by matching (or specific hybridization) or by recognition of an affinity site of the nucleic acids which are to be detected or identified in the biological sample.

Particular examples of the documents describing techniques relative to bioDNA chips are:

- the review article by Wang J. (Nucleic Acids Research, 28, 16:3011-3016, 2000), which has an abstract making the point on the main known techniques relative to DNA chips,
- the patent document issued under N° US 6,030,782, which describes grafting with a mercaptosilanized surface, of nucleic acids modified by a sulhydryl or disulfide group, and the article by Bamdad (Biophysical Journal, 75:1997-2003, 1998), which describes obtaining surfaces having DNA by incorporation of composite molecules, DNA-thiols, in auto-assembled monolayers (« self-assembled monolayers or SAMs »);
- the international patent application published under N° W0 00/43539 which proposes immobilizing molecules, such as oligonucleotides, by means of polyfunctional polymers (« polymer brushes ») thus enabling the grafting density to be increased. These polymers can be obtained from hydroxyethyl, acrylamide methacrylate, or vinyl pyrrolidone;
- the international patent application published under N° WO 00/36145 describes a fabrication method of DNA chips, comprising polymerization on a substrate of metallic layer type, a copolymer of pyrrol and functionalized pyrrol, fixing a reticulation agent on the functionalized pyrrol, then fixing a biological probe (such as an

oligonucleotide). The reticulation agent can be bifunctional, and for example have an ester function of the N-hydroxysuccinimide and a maleimide function;

- the international patent application published under N° WO 98/20020 which likewise describes the high-density immobilization of nucleic acids on solid supports, this time by placing in contact of a nucleic acid containing a thiol group with support having a group reacting with this thiol, possibly by way of a reticulation agent;

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- the article by Penchovsky *et al.* (Nucleic Acids Research, 28, 22, e98, 2000), which describes a method for immobilization of oligonucleotides on aminated balls of latex, by means of a reticulation agent which reacts under the action of light; and
- the international patent applications published under N° WO 99/16907, WO 00/40593 and WO 00/44939 filed by the company Surmodics (which produces lames for registering oligonucleotides functionalized with an amine). These applications describe especially the fixing of nucleic acids on surfaces such as glass, by way of a polymer skeleton to which one or more « photochemically active » groups are fixed on one side of the polymer (for grafting on the surface) and « thermochemically active » on the other side (for grafting with the functionalized nucleic acid).

A nucleotidic sequence (probe or primer) according to the invention thus enables detection and/or amplification of specific nucleic sequences. In particular, detection of said sequences is made easier when the probe is fixed on a DNA chip, or to a high-density filter.

The utilization of DNA chips or of high-density filters in effect helps determine expression of genes in an organism having a genomic sequence close to the genome of *Legionella pneumophila* Paris strain (Collection of the Pasteur Institute CIP 107-629-T).

The genomic sequence of *Legionella pneumophila* Paris strain, completed by identification of all the genes of this organism, such as presented in the present invention, serves as a base for the construction of these DNA chips or filter.

The preparation of these filters or chips consists of synthesizing oligonucleotides, corresponding to the 5' and 3' ends of the genes. These oligonucleotides are selected by utilizing the genomic sequence and its annotations divulged by the present invention. The matching temperature of these oligonucleotides at the corresponding places on the DNA must be approximately the same for each oligonucleotide. This aids in preparing fragments of DNA corresponding to each gene by the utilization of condition of appropriate PCR in a highly automated environment.

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The amplified fragments are then immobilized on filters or supports made of glass, silicon or synthetic polymers and these media are utilized for hybridization.

The availability of such filters and/or chips and of the annotated corresponding genomic sequence allows study of the expression of large ensembles, or of the totality of genes in the microorganisms of the *Legionella* genre or of the *Legionella pneumophila* species, by preparing the complementary DNA, and by hybridizing it to the DNA or to the oligonucleotides immobilized on the filters or the chips. Likewise, the filters and/or the chips allow study of the variability of the strains or species, by preparing the DNA of these organisms and by hybridizing it to the DNA or to the oligonucleotides immobilized on the filters or the chips.

The differences between the genomic sequences of the different strains or species can extensively affect the intensity of the hybridization and, as a consequence, perturb interpretation of the results. It can thus be necessary to have the precise sequence of the genes of the strain to be studied. The method for detecting the genes described in detail hereinbelow, implying determination of the sequence of random fragments of a genome, and organizing it according to the sequences of the complete genome of the *Legionella pneumophila* strain divulged in the present invention, can be very useful.

The nucleotide, or proteic, sequences according to the invention can be likewise utilized in DNA chips, or, if required, protein chips for performing mutation analysis. This analysis is based on the constitution of chips, especially DNA chips, capable of analyzing each base of a nucleotidic sequence according to the invention. For this purpose the techniques of micro-sequencing on DNA chip especially could be used. The mutations are detected by extension of immobilized primers hybridizing analyzed sequences to the matrix, just in a position adjacent to that of the desired mute nucleotide. A single-strand matrix, RNA or DNA, of the sequences for analysis will be advantageously prepared according to classic methods, from products amplified according to PCR-type techniques. The matrices of single-strand DNA, or RNA thus obtained are then deposited on the DNA chip, in conditions enabling their specific hybridization to the immobilized primers. A thermostable polymerase, for example Tth or the Taq DNA polymerase, specifically extends the end 3' of the primer immobilized with an analog of complementary marked nucleotide of the nucleotide in a variable position of the site; for example a thermal cycle is created in the presence of fluorescent dideoxyribonucleotides. The experimental conditions will be adapted especially to the

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chips employed, to the immobilized primers, to the polymerases employed, and to the marking system selected. An advantage of microsequencing, relative to techniques based on hybridization of probes, is that it allows all the variable nucleotides to be identified with optimal discrimination in conditions of homogeneous reactions; when used on DNA chips, it permits optimal resolution and specificity for routine and industrial detection of mutations in multiplex.

The utilization of high-density filters and/or chips thus provides new knowledge on the regulation of genes in organisms of industrial importance, and in particular the legionelloses propagated in diverse conditions. It also allows rapid identification of the differences between the genome of the strains utilized in multiples industrial applications.

In addition, a DNA chip or a filter can be an interesting extremely tool for determination, detection and/or identification of a microorganism. Therefore, the DNA chips according to the invention which further contain at least one nucleotidic sequence of a microorganism other than *Legionella pneumophila* Paris strain, immobilized on the support of said chip are likewise preferred. Preferably, the selected microorganism is selected from among the species of bacteria of the *Legionella* genre (hereinbelow designated at times as bacteria associated with *Legionella pneumophila*), or the subspecies of *Legionella pneumophila*, or again the variants of *Legionella pneumophila* Paris strain.

A DNA chip or a filter according to the invention is a very useful element of certain kits or is necessary for the detection and/or identification of microorganisms, in particular the bacteria belonging to the *Legionella pneumophila* species Paris strain or the associated microorganisms, likewise objects of the invention.

Besides, the DNA chips or the filters according to the invention, containing probes or specific primers of *Legionella pneumophila* Paris strain, compared in particular to the *Philadelphia* strain, are highly advantageous elements of kits or necessary for the detection and/or quantification of the expression of genes of *Legionella pneumophila* Paris strain (or of associated microorganisms).

In effect, control of the expression of genes is a critical point for optimizing the growth and yield of a strain, either allowing the expression of one or more novel genes, or by modifying the expression of genes already present in the cell. The present invention provides all the sequences naturally active in *Legionella pneumophila* Paris strain enabling the expression of the genes. It thus allows all the sequences expressed in

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Legionella pneumophila Paris strain to be determined. It likewise provides a tool for locating the genes whereof the expression follows a given pattern. To achieve this, the DNA of all or part of the genes of Legionella pneumophila Paris strain can be amplified thanks to primers according to the invention, then fixed to a support, such as for example glass or nylon or a DNA chip, so as to construct a tool allowing the expression profile of these genes to be followed. This tool, constituted by this support containing the coding sequences serves as hybridization matrix to a mixture of marked molecules reflecting the RNA messengers expressed in the cell (in particular the probes marked according to the invention). By repeating this experience at different instants and by combining all these data via suitable processing, the expression profiles of all these genes are obtained. The knowledge of the sequences which follow a given regulation pattern can also be of benefit for direct searching, for example by homology, for other sequences following globally, but slightly different for the same regulation pattern. By way of complement, it is possible to isolate each control sequence present upstream of the segments serving as probes and to follow their activity by means of appropriate means such as a reference gene (luciferase, β-galactosidase, GFP for «Green Fluorescent Protein »). These isolated sequences can then be modified and assembled by metabolic engineering with sequences of interest with a view to their optimal expression.

The aim of the invention likewise is the cloning and/or expression vectors, which contain a nucleotidic sequence according to the invention. In particular, those nucleotidic sequences are preferred which code for polypeptides having a cellular envelope or surface, or implied in the cellular machinery, in particular secretion, central intermediary metabolism, in particular production of sugar, energetic metabolism, the synthesis process of Vitamin B12, transcription and translation, synthesis of polypeptides.

The vectors according to the invention preferably comprise elements which permit the expression and/or secretion of the nucleotidic sequences in a determined host cell.

The vector must comprise a promoter, signals for initiation and termination of translation, thus of the appropriate regulation regions of transcription. It must be able to be kept stable in the host cell and can possibly have particular signals which specify secretion of the translated protein. These different elements are selected and optimized

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by the specialist as a function of the cellular host utilized. To this effect, the nucleotidic sequences according to the invention can be inserted into autonomous replication vectors within the selected host, or they can be integrative vectors of the selected host.

Such vectors are prepared by methods currently utilized by the specialist, and the resulting clones can be introduced to an appropriate host using standard methods, such as lipofection, electroporation, thermal choc, or chemical methods.

The vectors according to the invention are for example vectors of plasmidic or viral origin. They are useful for transforming host cells so as to clone or express the nucleotidic sequences according to the invention.

The invention likewise comprises the host cells transformed by a vector according to the invention.

The cellular host can be selected from amongst prokaryotic or eukaryotic systems, for example bacterial cells, but likewise yeast cells or animal cells, in particular the cells of mammals. The cells of insects or plant cells can likewise be used here. The host cells preferred according to the invention are in particular prokaryotic cells, preferably bacteria such as *E. coli*, or again belonging to the *Legionella* genre, to the *Legionella pneumophila* species Paris strain, or microorganisms associated with the *Legionella pneumophila* species Paris strain.

The invention relates likewise to animals, except human, which comprise a cell transformed according to the invention. The transformed cells according to the invention are utilizable in preparation processes for recombinant polypeptides according to the invention. The preparation processes of a polypeptide according to the invention in recombinant form, characterized in that they utilize a vector and/or a cell transformed by a vector according to the invention are themselves included in the present invention. Preferably, a cell transformed by a vector according to the invention is cultivated in conditions permitting expression of said polypeptide and said recombinant peptide is recovered. The host cells according to the invention can likewise be utilizes for preparation of nutritive compositions, themselves an object of the present invention.

As was mentioned, the cellular host can be selected from amongst prokaryotic or eukaryotic systems. In particular, it is possible to identify nucleotidic sequences according to the invention, facilitating secretion in such a prokaryotic or eukaryotic system. A vector according to the invention carrying such a sequence can thus be used advantageously for production of recombinant proteins, to be secreted. In fact,

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purification of these recombinant proteins of interest will be facilitated by the fact that they are present in the supernatant of the culture cellular rather than inside the host cells.

The polypeptides according to the invention can likewise be prepared by chemical synthesis. Such a preparation process is likewise an object of the invention. The specialist is aware of synthesis chemical processes, for example techniques utilizing solid phases (see especially Steward *et al.*, 1984, Solid phase peptides synthesis, Pierce Chem. Company, Rockford, 111, 2nd ed., (1984)) or techniques utilizing partial solid phases, by condensing fragments or by synthesis in classic solution. The polypeptides obtained by chemical synthesis and capable of comprising corresponding non-natural amino acids are likewise included in the invention.

The invention is further relative to hybrid polypeptides having at least one polypeptide or one of its fragments according to the invention, and a sequence of a polypeptide for inducing an immune response in humans or animals.

Advantageously, the antigenic determinant is such that it is capable of inducing a humoral and/or cellular response.

Such a determinant could comprise a polypeptide or one of its fragments according to the invention in glycosylated form utilized with a view to obtaining immunogenic compositions capable of inducing synthesis of antibodies directed against multiple epitopes. Said polypeptides or their glycosylated fragments likewise part of the invention.

These hybrid molecules can be made up in part by a carrier molecule of polypeptides or of their fragments according to the invention, associated with a possibly immunogenic part, in particular an epitope of the diphtheric toxin, tetanic toxin, a surface antigen of the hepatitis B virus (patent FR 79 21811), the antigen VP1 of the poliomyelitus virus or any other toxin or viral or bacterial antigen.

The synthesis processes of hybrid molecules encompass the methods utilized in genetics to construct hybrid nucleotidic sequences coding for the desired polypeptidic sequences. For example, reference can be made advantageously to the technique for obtaining genes coding for fusion proteins described by Minton in 1984.

Said hybrid nucleotidic sequences coding for a hybrid polypeptide, as well as the hybrid polypeptides according to the invention characterized in that these are recombinant polypeptides obtained by the expression of said hybrid nucleotidic sequences, are likewise part of the invention.

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The invention likewise comprises the vectors characterized in that they contain one of said hybrid nucleotidic sequences. The host cells transformed by said vectors, the transgenic animals comprising one of said transformed cells as well as the preparation processes for recombinant polypeptides utilizing said vectors, said transformed cells and/or said transgenic animals are likewise naturally part of the invention.

The coupling between a polypeptide according to the invention and an immunogenic polypeptide can be made chemically, or biologically. Therefore, according to the invention, it is possible to introduce one or more binding element(s), especially amino acids for facilitating the coupling reactions between the polypeptide according to the invention, and the immunostimulator polypeptide, the covalent coupling of the immunostimulator antigen able to be formed at the N or C-terminal end of the polypeptide according to the invention. The bifunctional reagents allowing this coupling are determined as a function of the end selected for making this coupling, and the coupling techniques are well known to the specialist.

The conjugates originating from a coupling of peptides can likewise be prepared by genetic recombination. The (conjugated) hybrid peptide can in effect be produced by recombinant DNA techniques, by insertion in or addition to the DNA sequence coding for the polypeptide according to the invention, of a sequence coding for the antigenic, immunogenic or haptenic peptide(s). These preparation techniques for hybrid peptides by genetic recombination are well known to the specialist (see for example Makrides, 1996, Microbiological Reviews 60:512-538).

Preferably, said immune polypeptide is selected in the group of peptides containing anatoxins, especially diphteric toxoid or tetanic toxoid, the proteins derived from Streptococcus (as the binding protein to human seralbumin), the membranous OMPA proteins and the complexes of proteins of external membranes, the vesicles of external membranes or the proteins of thermal shocks.

The hybrid polypeptides according to the invention are very useful to obtain monoclonal or polyclonal antibodies, capable of specifically recognizing the polypeptides according to the invention. In effect, a hybrid polypeptide according to the invention allows potentiation of the immune response, against the polypeptide according to the invention coupled to the immunogenic molecule. Such monoclonal or polyclonal antibodies, their fragments, or chimeric antibodies, recognizing the polypeptides according to the invention, are likewise objects of the invention.

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The specific monoclonal antibodies can be obtained according to the classic method of hybridome culture described by Köhler and Milstein (1975, Nature 256, 495).

The antibodies according to the invention are for example chimeric antibodies, humanized antibodies, Fab, or F(ab')² fragments. It can likewise be in the form of immunoconjugate or antibodies marked so as to provide a detectable and/or quantifiable signal.

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Therefore, the antibodies according to the invention can be employed in a process for detection and/or identification of bacteria belonging to the *Legionella pneumophila* species Paris and/or Lens strain and/or Philadelphia, or to an associated microorganism in a biological sample, characterized in that it comprises the following stages:

- a) contact of the biological sample with an antibody according to the invention;
- b) evidence of the possibly formed antigen-antibody complex.

The antibodies according to the present invention are likewise utilizable for detecting an expression of a gene of Legionella pneumophila Paris and/or Lens strain and/or Philadelphia strain, or of associated microorganisms. In effect, the presence of the expression product of a gene recognized by a specific antibody of said product expression can be detected by the presence of an antigen-antibody complex formed after contact of the strain of Legionella pneumophila Paris, Lens or Philadelphia strain, or of the associated microorganism with an antibody according to the invention. The bacterial strain utilized can were «prepared», that is, centrifuged, lysated, placed in an appropriate reagent for the constitution of the medium prone to immunological reaction. In particular, a detection process for expression in the gene is preferred, corresponding to a Western blot, capable of being carried out after electrophoresis on polyacrylamide gel of a lysate of the bacterial strain, in the presence or in the absence of reductory conditions (SDS-PAGE). After migration and separation of the proteins on the polyacrylamide gel, said proteins are transferred to an appropriate membrane (for example made of nylon) and the presence of the protein or of the polypeptide of interest is detected, by contact of said membrane with an antibody according to the invention.

Therefore, the present invention likewise comprises the kits or the necessary for implementing a process such as described (detection of the expression of a gene of *Legionella pneumophila* Paris and/or Lens strain and/or Philadelphia, or of an associated microorganism, or for detection and/or identification of bacteria belonging to

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the Legionella pneumophila species Paris and/or Lens and/or Philadelphia strain, or an associated microorganism), comprising the following elements:

- a) a polyclonal or monoclonal antibody according to the invention;
- b) possibly, the reagents for the constitution of the medium prone to immunological reaction;
- c) possibly, the reagents allowing use of the antigen-antibody complexes produced by the immunological reaction.

The polypeptides and the antibodies according to the invention can advantageously be immobilized on a support, especially a protein chip. This type of protein chip is an object of the invention, and can likewise contain at least one polypeptide of a microorganism other than *Legionella pneumophila* Paris and/or Lens strain and/or Philadelphia strain, or an antibody directed against a compound of a microorganism other than *Legionella pneumophila* Paris and/or Lens and/or Philadelphia strain.

The high-density protein or filter chips containing proteins according to the invention can be constructed in the same way as the DNA chips according to the invention. In practice, synthesis of the polypeptides fixed directly on the protein chip, or ex situ synthesis followed by a stage for fixing the synthesized polypeptide on said chip can be carried out. This latter method is preferable, when proteins of significant size are to be fixed on the support, which are advantageously prepared by genetic engineering. All the same, if it is preferred to fix only peptides on the support of said chip, it can be more interesting to proceed with synthesis of said peptides directly in situ.

The protein chips according to the invention can be advantageously utilized in kits or necessary for the detection and/or identification of bacteria associated with the Legionella pneumophila species Paris and/or Lens strain and/or Philadelphia strain, or with a microorganism, more generally in kits or necessary for the detection and/or identification of microorganisms. When the polypeptides according to the invention are fixed on DNA chips, the presence of antibodies is searched for in the samples tested, with the fixation of an antibody according to the invention on the support of the protein chip allowing identification of the protein whereof said antibody is specific.

Preferably, an antibody according to the invention is fixed on the support of the protein chip, and the presence of the corresponding antigen, specific to *Legionella pneumophila* Paris and/or Lens strain and/or Philadelphia strain or of an associated microorganism is detected.

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A protein chip described hereinabove can be utilized for the detection of products of genes, for establishing an expression profile of said genes, as a complement to a DNA chip according to the invention.

The protein chips according to the invention are likewise extremely useful for proteomic experiments, which studies the interactions between the different proteins of a given microorganism. In a simplified manner, peptides representative of the different proteins of an organism are fixed on a support. Next, said support is put in contact with marked proteins, and after an optional rinsing stage, interactions are detected between said marked proteins and the peptides fixed on the protein chip.

Therefore, the protein chips comprising a polypeptidic sequence according to the invention or an antibody according to the invention are an object of the invention, as well as kits or necessary containing them.

The present invention likewise covers a process for detection and/or identification of bacteria belonging to the *Legionella pneumophila* species Paris and/or Lens and/or Philadelphia strain, or to an associated microorganism in a biological sample, which uses a nucleotidic sequence according to the invention.

It should be understood that the term biological sample relates in the present invention to the samples taken from a living organism (in particular blood, tissue, organs or the like taken a mammal) or a sample containing biological material, that is, DNA. Such a biological sample especially includes all fluids (liquid or air-borne), or any object, such as conduits for fluid, filters for fluids, or any object capable of being implied in the fluid supply in buildings, nutritional compositions containing bacteria or other.

The process for detection and/or identification utilizing the nucleotidic sequences according to the invention can be diverse in nature.

A process comprising the following stages is preferred:

- a) possibly, isolation of DNA from the biological sample to be analyzed, or obtaining DNAc from the RNA of the biological sample;
- b) specific amplification of the DNA of bacteria belonging to the *Legionella pneumophila* species Paris strain, Lens, Philadelphia or to a microorganism associated by means of at least one primer according to the invention;
 - c) revealing amplification products.

This process is based on specific amplification of DNA, in particular by a chain amplification reaction.

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Likewise, a process comprising the following stages is preferred:

- a) contact by a nucleotidic probe according to the invention with a biological sample, the nucleic acid contained in the biological sample having, if required, previously been made accessible to hybridization, in conditions permitting hybridization of the probe to the nucleic acid of bacteria belonging to the *Legionella pneumophila* species Paris strain, or to an associated microorganism;
- b) revealing the hybrid possibly formed between the nucleotidic probe and the DNA of the biological sample.

Such a process must not be limited to detection of the presence of the DNA contained in the biological sample tested, and it can likewise be used for detecting the RNA contained in said sample. This process in particular includes the Southern and Northern blot.

Another preferred process according to the invention comprises the following stages:

- a) contact by a nucleotidic probe immobilized on a support according to the invention with a biological sample, the nucleic acid of the sample, having, if required, been previously made accessible to hybridization, in conditions allowing hybridization of the probe to the nucleic acid of bacteria belonging to the *Legionella pneumophila* species Paris strain or to an associated microorganism;
 - b) contact by the hybrid formed between the nucleotidic probe immobilized on a support and the nucleic acid contained in the biological sample, if required after elimination of the DNA from the biological sample not having hybridized with the probe, with a nucleotidic probe marked according to the invention;
 - c) revealing the novel hybrid formed at stage b).

This process is advantageously utilized with a DNA chip according to the invention, the desired nucleic acid hybridizing with a probe present on the surface of said chip, and being detected by utilization of a marked probe. This process is advantageously implemented by combining a previous amplification stage of DNA or of complementary DNA obtained possibly by inverse transcription, by means of primers according to the invention.

Therefore, the present invention likewise includes the kits or necessary for the detection and/or identification of bacteria belonging to the *Legionella pneumophila* species Paris and/or Lens and/or Philadelphia strain, or to an associated microorganism, characterized in that it comprises the following elements:

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- a) a nucleotidic probe according to the invention;
- b) possibly, the reagents necessary for using a hybridization reaction;
- c) possibly, at least one primer according to the invention as well as the reagents necessary for amplification reaction of the DNA.

Similarly, the present invention likewise includes the kits or necessary for the detection and/or identification of bacteria belonging to the *Legionella pneumophila* species Paris strain or to an encircled microorganism, characterized in that it comprises the following elements:

- a) a nucleotidic probe, known as capture probe, according to the invention;
- 10 b) an oligonucleotidic probe, known as revelation probe, according to the invention;
 - c) possibly, at least one primer according to the invention as well as the reagents necessary for amplification reaction of the DNA.

Finally, the kits or necessary for the detection and/or identification of bacteria belonging to the *Legionella pneumophila* species Paris and/or Lens and/or Philadelphia strain, or to an associated microorganism, characterized in that it comprises the following elements:

- a) at least one primer according to the invention;
- b) possibly, the reagents necessary for performing an amplification reaction 20 of DNA;
 - c) possibly, a compound enabling the sequence of the amplified fragment, more particularly an oligonucleotidic probe according to the invention, to be verified, are likewise objects of the present invention.

Preferably, said primers and/or probes and/or polypeptides and/or antibodies according to the present invention utilized in the processes and/or kits or necessary according to the present invention are selected from amongst the primers and/or probes and/or polypeptides and/or antibodies specific to the *Legionella pneumophila* species Paris and/or Lens and/or Philadelphia strain. In a preferred manner these elements are selected from amongst the nucleotidic sequences coding for a secreted protein, among the polypeptides secreted, or among the antibodies directed against exported polypeptides, such as those implied in the wall or the cellular envelope of *Legionella pneumophila* Paris and/or Lens and/or Philadelphia strain.

The object of the present invention is likewise the strains of Legionella pneumophila Paris or Lens strain, and/or associated microorganisms containing one or

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more mutation(s), at most less than 10 % mutation (or again less as cited for the modifications of polypeptides) in a nucleotidic sequence according to the invention, in particular an ORF sequence, or their regulatory elements (in particular promoters).

According to the present invention, the strains of Legionella pneumophila Paris or Lens strain having one or more mutation(s) in the nucleotidic sequences coding for polypeptides implied in the machine cellular, in particular secretion, central intermediary metabolism, energetic metabolism, the process of synthesizing of the amino acids, transcription and translation, synthesis of the polypeptides, are preferred.

Said mutations can lead to inactivation of the gene, or in particular when they are situated in the regulatory elements of said gene, at overexpression of the latter.

The invention relates further to utilizing a compound selection method capable of inhibiting the expression of genes implied in the biosynthesis of polysaccharides having a cellular envelope of bacteria of the *Legionella pneumophila* species Paris strain, characterized in that it comprises the following stages:

- a) contact by said compound with a bacteria of said Paris strain, said bacteria being in conditions and in medium appropriate to its culture;
- b) determination of the capacity of said compound to inhibit the expression of the genes coding for the proteins of SEQ ID Nos. 1126, 3218, 288, 632, 917, 1503, 1555, 1877, 1928, 1963, 2204, 2212, 2243, 2324, 2378, 2410, 2411;
- c) by means of a process according to the invention in which said antibody is directed specifically against a polypeptide implied in the biosynthesis of the polysaccharides, or by means of a process according to the invention in which the probes or primers are specific to a nucleic sequence coding for a polypeptide implied in the biosynthesis of the polysaccharides;
- d) selection of organic or inorganic compound capable of modulating, regulating, inducing or inhibiting the expression of genes, and/or of modifying the cellular replication of eukaryotic or prokaryotic cells or capable of inducing, inhibiting or aggravating the pathologies associated with infection by *Legionella pneumophila* Paris strain or one of its associated microorganisms.

The invention likewise comprises a method for selection of compounds capable of binding to a polypeptide or one of its fragments according to the invention, capable of binding to a nucleotidic sequence according to the invention, or capable of recognizing an inventive antibody, and/or capable of modulating, regulating, inducing or inhibiting the expression of genes, and/or modifying the growth or cellular

replication of eukaryotic or prokaryotic cells, or capable of inducing, inhibiting or aggravating in a animal or human organism the pathologies bound to infection by *Legionella pneumophila* Paris strain or Lens or Philadelphia strains, or one of its associated microorganisms, characterized in that it comprises the following stages:

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a) contact by said compound with said polypeptide, said nucleotidic sequence, with a cell transformed according to the invention and/or administration of said compound to an animal transformed according to the invention;

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b) determination of the capacity of said compound to be bound with said polypeptide or said nucleotidic sequence, or to modulate, regulate, induce or inhibit the expression of genes, or to modulate the growth or cellular replication, or induce, inhibit or aggravate in said transformed animal the pathologies bound to infection by Legionella pneumophila Paris strain or Lens or Philadelphia strains strain, or one of its associated microorganisms.

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The cells and/or the animals transformed according to the invention, could advantageously serve as a model and be utilized in processes for studying, identifying and/or selecting compounds capable of being responsible for pathologies induced or aggravated by Legionella pneumophila Paris strain or Lens or Philadelphia strains strain, or capable of preventing and/or treating these pathologies such as for example genital, ocular or systemic diseases, especially of the lymphatic system. In particular, the transformed host cells, especially the bacteria of the family of Legionellae whereof the transformation by a vector according to the invention can for example grow or inhibit its infectious capacity, or modulate the pathologies habitually induced or aggravated by the infection, could be utilized for infecting animals in which the appearance of pathologies will be followed. These animals not transformed, infected for example with transformed Legionellae bacteria, will be able to serve as a study model. In the same manner, the animals transformed according to the invention will be able to be utilized in selection processes for compounds capable of preventing and/or treating diseases due to Legionella. Said processes utilizing said transformed cells and/or transformed animals are part of the invention.

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The compounds capable of being selected can be organic compounds such as polypeptides or hydrates of carbon or any other already known organic or inorganic compounds, or novel organic compounds elaborated by molecular modeling techniques and obtained by chemical or biochemical synthesis, these techniques being known to the specialist.

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Said selected compounds will be able to be utilized for modulating the growth and/or cellular replication of *Legionella pneumophila* Paris and/or Lens and/or Philadelphia strain, or any other associated microorganism and thus to control infection by these microorganisms. Said compounds according to the invention will likewise be utilized for modulating the growth and/or cellular replication of all eukaryotic or prokaryotic cells, especially tumoral cells and infectious microorganisms, for which said compounds will prove to be active, the methods determining said modulations being well known to the specialist.

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Compound capable of modulating the growth of a microorganism is understood to mean any compound allowing to intervene, modify, limit and/or reduce the development, growth, rate of proliferation and/or the viability of said microorganism.

This modulation can be realized for example by an agent capable of binding to a protein and thus inhibit or potentialize its biological activity, or capable of binding to a membranous protein of the external surface of a microorganism and blocking penetration of said microorganism in the host cell or benefiting the action of the immune system of the infected organism directed against said microorganism. This modulation can likewise be realized by an agent capable of binding to a nucleotidic sequence of DNA or RNA of a microorganism and for example blocking the expression of a polypeptide whereof the biological or structural activity is necessary to the growth or to the reproduction of said microorganism.

For these screening methods, likewise associated microorganism in the present selection method is understood to mean any microorganism whereof the gene expression can be modulated, regulated, induced or inhibited, or whereof the growth or cellular replication can be likewise modulated by a compound of the invention. Likewise, associated microorganism in the present invention is understood to mean any microorganism comprising nucleotidic sequences or polypeptides according to the invention. These microorganisms can in certain cases comprise polypeptides, or nucleotidic sequences identical or homologous to those of the invention will likewise be able to be detected and/or identified by the processes or detection and/or identification kit according to the invention and likewise serve as target for the compounds of the invention.

The invention relates to the compounds capable of being selected by a selection method according to the invention.

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The invention likewise relates to a pharmaceutical composition comprising a compound selected from amongst the following compounds:

- a) a nucleotidic sequence according to the invention;
- b) a polypeptide according to the invention;
- c) a vector according to the invention;
- d) an antibody according to the invention; and
- e) a compound capable of being selected by a selection method according to the invention, possibly in association with a pharmaceutically acceptable vehicle.

Efficacious quantity is understood to mean an adequate quantity of said compound or antibodies, or of polypeptide of the invention, allowing to modulate the growth of *Legionella pneumophila* Paris and/or Lens and/or Philadelphia strain, or of an associated microorganism.

The invention also relates to a pharmaceutical composition according to the invention for the prevention or treatment of an infection by a bacteria belonging to the *Legionella pneumophila* species Paris strain or Lens or Philadelphia strains, or by an associated microorganism.

The further aim of the invention is an immunogenic and/or vaccinal composition, characterized in that it comprises one or more polypeptides according to the invention and/or one or more hybrid polypeptides according to the invention.

The invention also comprises utilization of a cell transformed according to the invention, for the preparation of a vaccinal composition.

The aim of the invention likewise is a vaccinal composition, characterized in that it contains a nucleotidic sequence according to the invention, a vector according to the invention and/or a cell transformed according to the invention.

The invention likewise relates to vaccinal compositions according to the invention, for the prevention or treatment of an infection by a bacteria belonging to the *Legionella pneumophila* species Paris strain or Lens or Philadelphia strains, or by an associated microorganism.

In a preferred manner the immunogenic and/or vaccinal compositions according to the invention for preventing and/or treating infection by *Legionella pneumophila* Paris strain or Lens or Philadelphia strains, or by an associated microorganism will be selected from among the immunogenic and/or vaccinal compositions comprising a polypeptide or one of its fragments corresponding to a protein, or one of its fragments, of the cellular envelope of *Legionella pneumophila* Paris strain or Lens or Philadelphia

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strains. The vaccinal compositions comprising nucleotidic sequences will preferably likewise comprise nucleotidic sequences coding for a polypeptide or one of its fragments corresponding to a protein, or one of its fragments, of the cellular envelope of *Legionella pneumophila* Paris strain or Lens or Philadelphia strains.

Of these preferred immunogenic and/or vaccinal compositions, the most preferred are those comprising a polypeptide or one of its fragments, or a nucleotidic sequence or one of its fragments whereof the sequences are selected from among the nucleotidic sequences or amino acids identified in this functional group and listed previously.

The polypeptides of the invention or their fragments entering the immunogenic compositions according to the invention can be selected by techniques known to the specialist such as for example on the capacity of said polypeptides to stimulate the T cells, which translates for example by their proliferation or the secretion of interleukins, and which terminates with the production of antibodies directed against said polypeptides.

In mice, in which a ponderal dose of the vaccinal composition comparable to the dose utilized in humans is administered, the reaction antibody is tested by taking serum followed by studying the formation of a complex between the antibodies present in the serum and the antigen of the vaccinal composition, according to customary techniques.

According to the present invention, said vaccinal compositions will preferably be in association with a pharmaceutically acceptable vehicle and, if required, with one or more adjuvants of appropriate immunity.

These days, diverse types of vaccines are available for protecting humans against infectious diseases: attenuated living microorganisms (M. bovis - BCG for tuberculosis), inactive microorganisms (flu virus), acellular extracts (Bordetella pertussis) for pertussis), recombinant proteins (surface antigen of the hepatitis B virus), polyosides (pneumococci). Vaccines prepared from synthesis peptides or genetically modified microorganisms expressing heterologous antigens are in experimentation. Still more recently, recombined plasmidic DNAs carrying coding for protector antigens were proposed as an alternative vaccinal strategy. This type of vaccination is realized with a particular plasmid deriving from a plasmid of E. coli which does not replicate in vivo and which codes solely for the vaccinating protein. Animals were immunized by simply injecting naked plasmidic DNA into the muscle. This technique results in expression of the vaccinal protein in situ and to an immune response of cellular type (CTL) and of

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humoral type (antibodies). This double induction of the immune response is one of the principal advantages of the vaccination technique with naked DNA.

The vaccinal compositions comprising nucleotidic sequences or vectors in which said sequences are inserted are described in particular in international application N° WO 90/11092 and likewise in international application N° WO 95/11307.

The nucleotidic sequence making up the vaccinal composition according to the invention can be injected into the host after having been coupled to compounds which benefit penetration of this polynucleotide inside the cell or its transport as far as the cellular nucleus. The resulting conjugates can be encapsulated in polymer microparticles, as described in international application N° WO 94/27238 (Medisorb Technologies International).

According to another embodiment of the vaccinal composition according to the invention, the nucleotidic sequence, preferably a DNA, is complexed with DEAE-dextran, with nuclear proteins, with lipids or encapsulated in liposomes or introduced in the form of a gel facilitating its transfection in the cells. The polynucleotide or the vector according to the invention can also be in suspension in a buffer solution or be associated with liposomes.

Advantageously, such a vaccine will be prepared according to the technique described by Tacson *et al.* or Huygen *et al.* in 1996 or again according to the technique described by Davis *et al.* in the international application N° WO 95/11307.

Such a vaccine can likewise be prepared in the form of a composition containing a vector according to the invention, placed under the control of regulation elements allowing its expression in humans or animals. For example, the polypeptidic antigen of interest, the pcDNA3 plasmid or the pcDNA1/neo plasmid could be utilized as an *in vivo* expression vector, both marketed by Invitrogen (R & D Systems, Abingdon, UK). Such a vaccine will comprise advantageously, apart from the recombinant vector, a saline solution, for example a sodium chloride solution.

Pharmaceutically acceptable vehicle is understood to mean a compound or a combination of compounds entering a pharmaceutical or vaccinal composition not causing secondary reactions and which enables for example ease of administration of active compound, an increase in its life expectancy and/or its efficacy in the organism, augmentation of its solubility in solution or again an improvement in its preservation. These pharmaceutically acceptable vehicles are well known and will be adapted by the

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specialist as a function of the nature and mode of administration of the selected active compound.

As for vaccinal formulations, these can comprise adjuvants of appropriate immunity which are known to the specialist, such as for example aluminum hydroxide, a representative of the family of muramyl peptides as one of the peptidic derivatives of N-acetyl-muramyl, a bacterial lysate, or even the incomplete Freund adjuvant.

Preferably, these compounds will be administered systemically, in particular intravenously, intramuscularly, intradermally or subcutaneously, or orally. In a more preferred way, the vaccinal composition comprising polypeptides according to the invention will be administered in several doses, spread out over time, intradermically or subcutaneously.

Their administration methods, posologies and optimal galenic forms can be determined according to the criteria generally taken into account in setting up treatment adapted to a patient such as for example age or body weight of the patient, the seriousness of the general status, tolerance to treatment and the secondary effects.

The invention comprises utilizing an inventive composition, for the treatment or prevention of diseases brought on or aggravated by the presence of *Legionella pneumophila* Paris strain or Lens or Philadelphia strains.

The invention comprises the utilization of a composition according to the invention for the treatment or prevention of systemic diseases, induced or aggravated by the presence of *Legionella pneumophila* Paris strain or Lens or Philadelphia strains.

Additionally, an object of the present invention likewise is a genomic DNA bank of a bacteria of the species *Legionella pneumophila* Paris strain, characterized in that this is the bank deposited with the CNCM on November 19, 2003, under the order number I-3138.

Additionally, an object of the present invention likewise is a genomic DNA bank of a bacteria of the species *Legionella pneumophila* Lens strain, characterized in that this is the bank deposited with the CNCM on September 23, 2004, under the order number I-3306.

Additionally, an object of the present invention likewise is a vector or a host cell as claimed in Claim 38 or 42, characterized in that this is the vector or the cell deposited with the CNCM on November 19, 2003, under the order number I-3137.

One of the advantages of using the BAC system relative to a cosmides system is that the plasmid utilized is present only in a maximum two copies per transformed cell,

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which reduces the potential for recombination between DNA fragments, and more importantly, which eliminates the risk of lethal overexpression of bacterial cloned genes. Nevertheless, the presence of BAC as a single copy signifies that the plasmidic DNA must be extracted from a large volume of culture in order to obtain enough DNA for the sequence. In addition, the stability and fidelity of maintenance of the clones in a BAC bank enable identification of genomic differences among different strains of *Legionella*, and identification of these genetic differences which can be responsible for the phenotypical variations observed between the different strains.

The genomic DNA banks described in the present invention effectively cover the genome of *Legionella pneumophila* Paris and Lens strains. All the same, although it is possible that certain regions have not been able to be cloned in said bank, by virtue of lethality problems in *Escherichia coli*, these regions can easily be amplified and identified by the specialist, by utilizing oligonucleotides specific to the sequences of the ends of the different clones which form the contigs.

Additionally, an object of the present invention likewise is a method for isolating a polynucleotide of interest present in a bacteria of the *Legionella* genre and absent from a bacteria of another genre, or present in a pathogenic bacteria of the *Legionella* genre, or again present in a bacteria of the *Legionella* pneumophila species and absent from a bacteria of any other species of the *Legionella* genre, or again present in a bacteria of the *Legionella* genre, or again present in a bacteria of the *Legionella* pneumophila species Paris and/or Lens and/or Philadelphia strain and absent from a bacteria of the *Legionella* pneumophila species of any other strain, characterized in that it utilizes at least the BAC bank deposited on November 19, 2003 (I-3138) with the C.N.C.M and the BAC bank deposited on September 23, 2004 (I-3306) with the CNCM according to the invention.

Said method is preferably characterized in that it comprises the following stages:

- a) isolating at least one polynucleotide contained in a clone of said DNA bank deposited with the CNCM on November 19 2003, under the order number I-3138 or contained in a clone of said BAC bank deposited on September 23 2004 under the number I-3306;
 - b) isolating:
- at least one genomic polynucleotide or DNAc of a second bacteria of another genre or of the *Legionella* genre, said second bacteria of the *Legionella* genre belonging to a different strain of the Paris strain or, alternatively,

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- at least one polynucleotide contained in a clone of a DNA bank based on a BAC prepared from the genome of a second bacteria of another genre or of the *Legionella* genre, said second bacteria of the *Legionella* genre belonging to a different strain of the Paris and/or Lens and/or Philadelphia strain;
- 5 c) hybridizing the polynucleotide of stage a) to the polynucleotide of stage b);
 - d) selecting the polynucleotides of stage a) which do not have the hybridization complex form with the polynucleotides of stage b); and
 - e) characterizing the selected polynucleotide.

The polynucleotide of stage a) can be prepared by the digestion of at least one recombinant BAC clone with an appropriate restriction enzyme, and optionally, amplification of the resulting polynucleotide insert.

Therefore, the method of the invention enables the specialist to effect comparative genomic studies between the different strains or species of the Legionella genre, for example between the pathogenic strains and their non-pathogenic equivalent.

In particular, it is possible to study and determine the regions of polymorphism between said strains.

LEGENDS OF THE FIGURES

Figure 1: Circular genomic map of the line L. pneumophila Paris and specific genes of the L. pneumophila Lens line. From the exterior: circle 1: genes of Paris line on the chains + and – respectively. Red line, inversion in line Lens. Color code: green, genes of Paris line, black, rRNA operons, red, known virulence genes; the numbers indicate their position: 1 lvh-lvr secretion system type IV (lvrABC, lvhB2B3B4B5, lvrD, lvhB6B8B9B10B11D4, lvrE); 2 dot/icm secretion system type IV (icmTSRQOMLKEGCDJBF); 3 mip, 4 lspA, 5 lspDE, 6 htrA, 7 lspFGHIJK, 8 enhABC, 9 dot/icm secretion system type IV (icmVWX and dotABCD), 10 momp; circle 2: specific genes of the Lens line relative to the Paris line; 3: bias G/C (G+C/G-C) of the Paris line; circle 4: G+C content of the Paris line with <32,5% G+C in light yellow, between 32.5% and 44.1% in yellow and with >44.1% G+C in dark yellow. The scale (Mb) is indicated on the outside, the origin of the replication in position 0.

Figure 2: Phylogenetic tree of a multiple sequential comparison of kinase domains from Legionella pneumophila of Paris line to other prokaryotic and eukaryotic kinases by utilizing the MEGA program. The calculation was made by utilizing the

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Poisson correction as a distance process and as a tree construction process the Neighbor-joining distance method. 0.2 indicates the 2 amino acid substitutions for 10 sites. The Nrprot access numbers, the names of genes and the names of organisms are indicated on the pattern. The numbers indicate the priming values.

Figure 3: Schema representative the genome core and the sole gene complement of L. pneumophila Paris, Lens and Philadelphia lines. The orthologous genes were defined by the most adequate reciprocated FASTA comparisons. The threshold was defined at a maximum of 80 % of sequence identities and at a length ratio of 0.75 to 1.33. The coding sequences of Philadelphia were determined by the Genmark predictions utilizing the « CAAT-box » program and the sequence obtained on the site http://www.genome3.cpmc.columbia.edu/-legion/project/ (latest version).

Figure 4: A. Comparison of the protein-coding RTX genes of L. pneumophila AA100, Paris and Lens lines. The sequence of the rtxA locus of line AA100 was obtained from the NCBI database (AAD41583). The dotted lines indicate that the correct number of repetitions is uncertain. B. Consensus sequences of the highly preserved repeated patterns of Paris and Lens lines. The amino-acid sequences in black indicate 11 amino-acids of the preserved N-terminal sequence of Paris and Lens lines, the amino-acids sequences in color represent the repeated patterns of each line (same color as for A). The underlined amino-acids indicate the positions which can change among the repetitions.

Figure 5: Pattern illustration of the different stages of intracellular growth of L. pneumophila in the macrophages. The different phases are numbered 1) Adhesion and invasion of L. pneumophila in the host cell 2) The phagosome does not fuse with the lysosomes but recruits organelles and converts to a compartment of rugged endoplasmic reticulum type. 3) Intracellular replication, non-flagellated L. pneumophila inside a phagosome. 4) Release of L. pneumophila. Flagellated. In red: Different important stages in the infectious cycle of L. pneumophila. In blue: Hypothesis indicating the stages at which the identified proteins could interfere in this cycle.

Figures 6 and 7: Southern Blot showing the specificity of the repeated sequence SEQ ID N° 7074 in L. pneumophila; the legend of Figure 6 given by Table XXV and that of Figure 7 by Table XXVI.

EXAMPLES

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Example 1: Materials and methods

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1. Construction of banks

Shotgun bank of small fragments (size 1.5 to 2.5 kb)

The chromosomal DNA of the strains studied was prepared by a classic method including proteinase K treatment and phenol extraction (9). Approximately 36 ug of DNA were broken by nebulization (1 minute under pressure of 1 bar) (4). The ends of the DNA fragments were rendered free by having the DNA-polymerase of the bacteriophage T4 act for 15 minutes at 37°C in the presence of 4 tri-phosphate nucleotides. The enzyme was inactivated by incubation of 15 mn at 75°C. Adaptors (invitrogen Cat. N° 408-18) were ligatured to these ends. After ligature, the fragments of chromosomal DNA of a size between 1500 and 2500 base pairs were purified after electrophoresis on agarose gel. The vector utilized for construction of the bank, pcDNA2.1 (Invitrogen), was digested by the BstX1 enzyme and purified by geneclean (BIO-101) after electrophoresis on agarose gel. The chromosomal DNA and the purified vector were ligatured by action of the ligase of the bacteriophage T4. The ligation mixture was introduced by transformation to the strain of *Escherichia coli* XL2-blue (Stratagene). Environ 4000 colonies are obtained per ul of the ligation mixture.

Bank of average fragments (size 5 to 10 kb)

The bank was constructed by the technique of 'partial fill in' in the vector pSYX34 (12). The chromosomal DNA of the strain *L. pneumophila* Paris was prepared by partial digestion by *Sau*IIIA (Roche). After precipitation of the DNA in sodium acetate and the stage of partial fill-in with the A and G nucleotides by utilizing the Klenow enzyme, the fragments of chromosomal DNA having a size of between 5000 and 10000 base pairs were purified after electrophoresis on agarose gel and geneclean.

The vector is prepared in the same way by partial digestion with the SalI enzyme, precipitation in sodium acetate then reaction of partial fill-in with the C and T nucleotides and purification on agarose gel and geneclean. The fragments of chromosomal DNA and the purified vector were ligatured by action of the ligase of the bacteriophage T4. The ligation mixture was introduced by transformation to the strain of Escherichia coli XL10-Gold (Stratagene). Around 4000 colonies are obtained per ul of the ligation mixture. The two ends of around 4000 fragments of this bank were sequenced.

Bank of large fragments (size 25 to 90 kb)

The bank of large fragments was constructed as described previously (4) by utilizing the pIndigo BAC vector (Epicentre). Briefly, in order to avoid mechanical

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breaking of the DNA molecules the cells were included in agarose blocks in which DNA extraction is performed directly. For the preparation of large-sized fragments we performed partial digestion by *Hind*III (Roche) and separation by electrophoresis in pulsed fields. Fragments of sizes between 40 and 80 and between 80 and 130 kb were excised from the gel, purified by agarase treatment and ligature with the vector. The ligation mixture was introduced by electroporation to the strain of *Escherichia coli* DH10B (Gibco BRL). 1300 colonies were stored. The plasmidic DNAs of these 1300 colonies were extracted and the two ends of the cloned fragments were sequenced.

2. Preparation of plasmids and sequencing.

The plasmids were prepared by a semi-automatic preparation method developed at the GMP laboratory and based on the alkaline lysis method (2). The chromosomal inserts were sequenced from their two ends by utilizing the T7 and universal primer in following the recommendations of the supplier (Applied-Biosystems). The sequences were determined by utilizing automatic sequencers of type 3700 (Applied-Biosystem).

3. Assembling of sequences.

The sequences were assembled by utilizing the software suite developed at the University of Washington, Phred, Phrap and Consed (5, 8). The sequence completion was done by utilizing the software suite CAAT-box (7). The finishing stage corresponds to resequencing of the regions where the sequence is only slightly secure and sequencing of the regions located between the contigs. It was carried out either by sequencing PCR products or by operating on the bank clones. The sequences of oligonucleotides were defined by utilizing the consed and Primo software (8, 10).

4. Annotation of sequences.

The identification of the phases coding (CDS) was done by utilizing the software suite CAAT-box (7). This program combines the results of different methods:

- (i) identification of open reading phases and their tri as a function of their size;
- (ii) analysis of the probability of being coding by utilizing the Genemark software (11);
- (iii) identification of translation start (initiation codon and fixing sequence of the ribosome); and
 - (iv) the % of identity of the proteic sequence deduced with the proteic sequences contained in sequence banks by utilizing BLASTP software.

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The functions of the proteins coded by the identified coding phases were predicted by analysis of the search results of similarities in the non-redundant NCBI bank (http://www.ncbi.nlm.nih.gov/BLAST/) by utilizing BLASTP software (1).

5. Comparison of the genomes – identification of the CDS specific to the strain of *L. pneumophila* Paris strain.

All the proteic sequences deduced from the predicted coding phases of each genome was compared to all the proteic sequences possibly coded by the other genome by using BLASTP software. A threshold of 75 % of identity on the totality of the length of the protein was retained to identify the specific proteins of an isolate. This very high value was retained since it best allows discrimination of the orthologous genes from the paralogous genes (6). For the proteic sequences for which the sequence preservation is high (> at 70 %) the preservation of the nucleotidic sequences of the genes will also be high and could give a signal in low-stringency hybridization conditions. It will be necessary to consider this eventuality in the analysis of the test result.

15 Example 2: Deposit of biological material

The following organisms were deposited on November 19, 2003 with the Collection Nationale de Cultures de Microorganisms (CNCM) [National Collection of Microorganism Cultures], 25 rue du Docteur Roux, 75724 Paris Cedex 15, France, according to the dispositions of the Budapest Treaty:

- Clone of a shotgun bank, clone in the pCDNA vector, of the genome of Legionella pneumophila Paris strain (Pasteur Institute Collection CIP 107-629-T), registered under the file number I-3137. The insert of this clone is at a size of 14.2 kb and contains a gene coding for an autotransporter called led0019A07;
 - BAC DNA bank (1248 clones) of the genome of *Legionella pneumophila* Paris strain (Pasteur Institute Collection CIP 107-629-T), registered under file number I-3138. Said bank BAC (I-3138) was made in the *E.coli* DH10B strain (Grant *et al.*, PNAS, 87:4645, 1990). The inserts of this bank were cloned in the pBelo BAC-Kan vector (Mozo *et al.*, Mol. Gen. Genet., 1998, 258:562-70) and have an average size of between 1.5 and 2.5 kb. The total of these inserts corresponds to complete coverage of the genome.

Example 3: Annotations of sequences

1. Genes specific to L. pneumophila Paris strain relative to the L. pneumophila Philadelphia strain

No significant identity between the nucleotidic sequence of the gene of L. pneumophila Paris strain and the genome of L. pneumophila Philadelphia strain.

Table VII: Example of annotation of sequences in the case of proteic and nucleic sequence of L. pneumophila Paris strain not having % of significant identity with respectively proteic and nucleic sequences of L. pneumophila Philadelphia strain

IPF No. of the gene of L. pneumophila Paris strain	IPF No. of the gene of L. pneumophila Philadelphia strain (best score)	% of identity of proteic sequences % of identity of nucleotic sequences	
2043.1	-	_	-
2094.2	-	-	-
2039.1	-	-	-
2051.2	3061.1	33 %	not significant
3425.1	5305.1	32 %	not significant

2. Genes common to the two strains *L. pneumophila* Paris strain and Philadelphia strain for which the % of identity of deduced nucleic and proteic sequences is less than 75 %

<u>Table VIII</u>: Example of annotation of sequences in the case of proteic and nucleic sequences of genes common to the two strains of *L. pneumophila* Paris strain and Philadelphia strain for which the % of identity of the deduced nucleic and proteic sequences is less than 75 %

IPF No. of the gene of L. pneumophila Paris strain	IPF No. of the gene of L. pneumophila Philadelphia strain (best score)	% of identity of proteic sequences	% of identity of nucleotidic sequences	
2244.2	3793.1	63 %	59 %	
258.2	1342.1	60 %	59 %	

3. Genes common to *L. pneumophila* Paris strain and Philadelphia strain for which the % of identity of the deduced nucleic and proteic sequences is greater than 75 %

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<u>Table IX</u>: Example of annotation of sequences in the case of proteic and nucleic sequences of genes common to the two strains of *L. pneumophila* Paris strain and Philadelphia strain for which the % of identity of the deduced nucleic and proteic sequences is greater than 75 %

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L.	IPF No. of pneumophila aris strain gene	IPF No. of L. pneumophila Philadelphia strain gene (best score)	% of identity of proteic sequences	% of identity of nucleotidic sequences
	4629.2	133.1	100 %	100 %
	6079.1	4147.1	90 %	88 %

Example 4: Example of alignment of sequences

Presented hereinbelow are the alignments of sequences preserved in the Paris and Philadelphia strains. For each of the six examples which follow, we present an alignment of the nucleotidic sequences as well as alignment of the sequences of amino acids. The alignment of the sequences of amino acids is obtained by aligning the translated sequence of the ORF present in the Paris strain with the sequence originating from translation in the six phases of the contigs of the Philadelphia sequence. The sequence homology of these ORFs present in the two strains is very strong, as much in amino acids as in nucleotides.

TBLASTN 2.2.6 [Apr-09-2003] Query= 1764.3 CONTIG=Contig42 POSCDS1=13736 POSCDS2=14869 SENS=p, Seq Id: 555 (216 letters) Database: contigsLpPhiladelphia 51 sequences; 3,410,887 total letters Searching.done Score E Sequences producing significant alignments: (bits) Value LpPhiladelphia Contig49 433 e-123 >LpPhiladelphia Contig49 Length = 376826Score = $433 \text{ bits (1114), Expect = } e^{-123}$ Identities = 215/215 (100%), Positives = 215/215 (100%) Frame = -2Ouerv: 1 HLALFDETYIKTILILLSICVLKKFILRYDMILNDIVLYDNFFMTFDYKDNFMSKGPYQS 60 HLALFDETYIKTILILLSICVLKKFILRYDMILNDIVLYDNFFMTFDYKDNFMSKGPYQS Sbjct: 297565 HLALFDETYIKTILILLSICVLKKFILRYDMILNDIVLYDNFFMTFDYKDNFMSKGPYQS 297386 Query: 61 FANRLISALKDRGYTASRSPNGICIKTLAEFTGASEQICRRYIRGDALPDYEKVKQLAFH ${\tt FANRLISALKDRGYTASRSPNGICIKTLAEFTGASEQICRRYIRGDALPDYEKVKQLAFH}$ Sbjct: 297385 FANRLISALKDRGYTASRSPNGICIKTLAEFTGASEQICRRYIRGDALPDYEKVKQLAFH 297206 LQVNPGWLLFGEDENATTKKNEVDEKLLHYILKOSHHLYPISOGSNDDYADFVLGLIKEV Query: 121 180 LQVNPGWLLFGEDENATTKKNEVDEKLLHYILKQSHHLYPISQGSNDDYADFVLGLIKEV Sbjct: 297205 LQVNPGWLLFGEDENATTKKNEVDEKLLHYILKQSHHLYPISQGSNDDYADFVLGLIKEV 297026 KAIDTSENNLLKIIDLAIGSISSYEEKRKKHSHAV KAIDTSENNLLKIIDLAIGSISSYEEKRKKHSHAV Sbjct: 297025 KAIDTSENNLLKIIDLAIGSISSYEEKRKKHSHAV 296921 Database: contigsLpPhiladelphia Posted date: Nov 20, 2003 10:38 AM Number of letters in database: 3,410,887 Number of sequences in database: 51 Lambda K 0.321 0.139 0.402 Gapped Lambda 0.267 0.0410 0.140 Matrix: BLOSUM62 Query= 1764.3 CONTIG=Contig42 POSCDS1=13736 POSCDS2=14869 SENS=p (1134 letters) Database: /home/Gmp/rusniok/projets/legionella/pourBrevet-191103/contigsLpPhiladelphia 51 sequences; 3,410,887 total letters Searching......done Score E

(bits) Value

Sequences producing significant alignments:

LpPhiladelphia Contig35 2248 0.0 >LpPhiladelphia Contig35 Length = 48622Score = 2248 bits (1134), Expect = 0.0Identities = 1134/1134 (100%) Strand = Plus / Plus Query: 1 atgatcagaaaaataatttatgttacaggtactcgtgccgattatggactgatgagagaa 60 Sbjct: 7091 atgatcagaaaaataatttatgttacaggtactcgtgccgattatggactgatgagagaa 7150 Query: 61 gtactaaaaagattacaccagtcagaagacattgacttatcgatttgtgtcactggtatg Sbjct: 7151 gtactaaaaagattacaccagtcagaagacattgacttatcgatttgtgtcactggtatg 7210 $\verb|catcttgatgctttgtatggaaatacagttaacgaaattaaagcagatcagttctcaata|\\$ Query: 121 Sbjct: 7211 catcttgatgctttgtatggaaatacagttaacgaaattaaagcagatcagttctcaata 7270 Query: 181 tgcggcattattcctgttgatcttgccaatgctcagcatagttctatggcaaaagctatc 240 Sbjct: 7271 tgcggcattattcctgttgatcttgccaatgctcagcatagttctatggcaaaagctatc 7330 Query: 241 ggccatgaacttttgggattcaccgaggtattcgaaagtgaaactcctgatgtcgtttta 300 Sbjct: 7331 ggccatgaacttttgggattcaccgaggtattcgaaagtgaaactcctgatgtcgtttta 7390 ttgctgggagatcgaggagaaatgcttgctgcggccatagcagcgatacatttaaatatc Query: 301 360 Sbjct: 7391 ttgctgggagatcgaggagaaatgcttgctgcggccatagcagcgatacatttaaatatc 7450 Query: 361 ccggttgtacatctgcacggaggagagcgctctggaaccgttgatgaaatggtaaggcat 420 Sbjct: 7451 ccggttgtacatctgcacggaggagagcgctctggaaccgttgatgaaatggtaaggcat 7510 gcgatttccaaattatctcattatcattttgtcgcaacagaggcatccaaacaacgattg Query: 421 Sbjct: 7511 gcgatttccaaattatctcattatcattttgtcgcaacagaggcatccaaacaacgattg 7570 Query: 481 attagaatgggtgagaaagaagaaaccatttttcaggttggtgctccaggcttggatgaa 540 Sbjct: 7571 attagaatgggtgagaaagaagaaccatttttcaggttggtgctccaggcttggatgaa 7630 atcatgcagtataaaacgtctacacgtgatgtgtttaatcaacgttatggatttgatcct 600 Query: 541 Sbjct: 7631 atcatgcagtataaaacgtctacacgtgatgtgtttaatcaacgttatggatttgatcct 7690 Query: 601 gacaaaaaaatctgtttattaatctatcacccggttgttcaagaagttgactcgattaaa

Sbjct: 7691 gacaaaaaaatctgtttattaatctatcacccggttgttcaagaagttgactcgattaaa 7750

Query: 661 attcaatttcaaagcgtgattcaggcagcactcgctacaaatttacagattatttgcctt 720
Query: 721 gagcctaattccgatacgggtggtcatttaattcgagaagtgattcaggaatatattgat 780
Query: 781 catcctgatgttagaattatcaagcacttacatcgtccggaatttattgattg
Query: 841 aattctgatgtgatgctgggaaattccagtagtggcatcatagaggcagcctcatttaac 900
Query: 901 ctgaacgtagttaatgttggaagcaggcaaaatttaagagaacgaagcgacaatgtcatt 960
Query: 961 gatgttgatgttacttatgatgctattttgactggtctaagagaagcgctaaataaa
Query: 1021 aagataaaatactctaactgttatggggatggaaaaacgagtgaaaggtgttatcaattg 1080
Query: 1081 ttaaaaactatccctttgcactcacaaatattgaataaatgcaatgcatactaa 1134 113
TBLASTN 2.2.6 [Apr-09-2003]
Query= 1864.3 CONTIG=Contig42 POSCDS1=77740 POSCDS2=79155 SENS=p, Seq Id: 622 (489 letters)
Database: contigsLpPhiladelphia 51 sequences; 3,410,887 total letters
Searching.done
Sequences producing significant alignments: Score E (bits) Value
LpPhiladelphia_Contig49 1003 0.
>LpPhiladelphia_Contig49 Length = 376826
Score = 1003 bits (2594), Expect = 0.0 Identities = 488/488 (100%), Positives = 488/488 (100%)

Frame = +2	
Query: 1 KLSLPLIRLWQLSRSKHMFKPQGLYDYICQQWQEEILPSLCDYIKIPNKSPHFDAKWEEH KLSLPLIRLWQLSRSKHMFKPQGLYDYICQQWQEEILPSLCDYIKIPNKSPHFDAKWEEH	60 ·
Sbjct: 21029 KLSLPLIRLWQLSRSKHMFKPQGLYDYICQQWQEEILPSLCDYIKIPNKSPHFDAKWEEH 21	208
GYMEQAVNHIANWCKSHAPKGMTLEIVRLKNRTPLLFMEIPGQIDDTVLLYGHLDKQPEM	120
Sbjct: 21209 GYMEQAVNHIANWCKSHAPKGMTLEIVRLKNRTPLLFMEIPGQIDDTVLLYGHLDKQPEM 21	388
SGWSDDLHPWKPVLKNGLLYGRGGADDGYSAYASLTAIRALEQQGLPYPRCILIIEACEE	180
Sbjct: 21389 SGWSDDLHPWKPVLKNGLLYGRGGADDGYSAYASLTAIRALEQQGLPYPRCILIIEACEE 21	568
SGSYDLPFYIELLKERIGKPSLVICLDSGAGNYEQLWMTTSLRGNLVGKLTVELINEGVH	240
Sbjct: 21569 SGSYDLPFYIELLKERIGKPSLVICLDSGAGNYEQLWMTTSLRGNLVGKLTVELINEGVH 21	748
SGSASGIVADSFRVARQLISRIEDENTGEIKLPQLYCDIPDERIKQAKQCAEILGEQVYS	300
Sbjct: 21749 SGSASGIVADSFRVARQLISRIEDENTGEIKLPQLYCDIPDERIKQAKQCAEILGEQVYS 21	928
EFPWIDSAKPVIQDKQQLILNRTWRPALTVTGADGFPAIADAGNVMRPVTSLKLSMRLPP	360
Sbjct: 21929 EFPWIDSAKPVIQDKQQLILNRTWRPALTVTGADGFPAIADAGNVMRPVTSLKLSMRLPP 22	108
Query: 361 LVDPEAASVAMEKALTQNPPYNAKVDFKIQNGGSKGWNAPLLSDWLAKAASEASMTYYDK LVDPEAASVAMEKALTQNPPYNAKVDFKIQNGGSKGWNAPLLSDWLAKAASEASMTYYDK	420
Sbjct: 22109 LVDPEAASVAMEKALTQNPPYNAKVDFKIQNGGSKGWNAPLLSDWLAKAASEASMTYYDK 22	288
Query: 421 PAAYMGEGGTIPFMSMLGEQFPKAQFMITGVLGPHSNAHGPNEFLHLDMVKKLTSCVSYV PAAYMGEGGTIPFMSMLGEQFPKAQFMITGVLGPHSNAHGPNEFLHLDMVKKLTSCVSYV	480
Sbjct: 22289 PAAYMGEGGTIPFMSMLGEQFPKAQFMITGVLGPHSNAHGPNEFLHLDMVKKLTSCVSYV 22	468
Query: 481 LYSFSQKK 488 LYSFSQKK	
Sbjct: 22469 LYSFSQKK 22492	
Database: contigsLpPhiladelphia Posted date: Nov 20, 2003 10:38 AM Number of letters in database: 3,410,887 Number of sequences in database: 51	
Lambda K H 0.318 0.136 0.419	
Gapped Lambda K H 0.267 0.0410 0.140	
Matrix: BLOSUM62	
Query= 1864.3 CONTIG=Contig42 POSCDS1=77740 POSCDS2=79155 SENS=p (1416 letters)	
Database: /home/Gmp/rusniok/projets/legionella/pourBrevet- 191103/contigsLpPhiladelphia 51 sequences; 3,410,887 total letters	
Searchingdone	
Score Sequences producing significant alignments: (bits) V	E alue

Ouery: 601

2807 0.0 LpPhiladelphia Contig49 >LpPhiladelphia_Contig49 Length = 376826Score = 2807 bits (1416), Expect = 0.0Identities = 1416/1416 (100%) Strand = Plus / Plus $\verb|atgttcaaaccccaaggattgtatgattacatatgccaacagtggcaagaagagatattg|$ 60 Query: 1 Sbjct: 21080 atgttcaaaccccaaggattgtatgattacatatgccaacagtggcaagaagagatattg 21139 ccaaqtttatgtgactacataaaaatccctaataaatctcctcactttgatgcaaaatgg 120 Query: 61 areimminimmuuuuuuuuuuuuuuuuuuuuu Sbjct: 21140 ccaagtttatgtgactacataaaaatccctaataaatctcctcactttgatgcaaaatgg 21199 qaaqaacatqqttatatqqaqcaqqcaqttaatcacattqccaattqgtqtaaqtcqcat 180 Query: 121 Sbjct: 21200 gaagaacatggttatatggagcaggcagttaatcacattgccaattggtgtaagtcgcat 21259 gctcccaaaggaatgactctggaaattgttcgcctgaaaaataggactccattactattt 240 Query: 181 Sbjct: 21260 gctcccaaaggaatgactctggaaattgttcgcctgaaaaataggactccattactattt 21319 atggaaattccaggccaaattgatgacactgtgttgctttatgggcacttggataaacaa 300 Query: 241 Sbjct: 21320 atggaaattccaggccaaattgatgacactgtgttgctttatgggcacttggataaacaa 21379 360 cctgagatgtcaggctggagtgacgatttacatccatggaaacccgtattgaaaaatgga Query: 301 mainiminiiriinmmmmimmimatuutuuti Sbjct: 21380 cctgagatgtcaggctggagtgacgatttacatccatggaaacccgtattgaaaaatgga 21439 420 Query: 361 amanninininininininininina amananinin Sbict: 21440 ttgttatacqqaaqaqqqqcaqatqatqqtattctgcttatgcatcactcacggct 21499 attcqcqccttqqaacaqcaaqqtttqccatatcctcqttqtatattaatcatcqaaqcq 480 Query: 421 Sbjct: 21500 attcgcgccttggaacagcaaggtttgccatatcctcgttgtatattaatcatcgaagcg 21559 tgtgaggaaagtggcagttacgatttgcctttttatattgagttgctgaaagagcgtatt 540 Query: 481 Sbjct: 21560 tgtgaggaaagtggcagttacgatttgcctttttatattgagttgctgaaagagcgtatt 21619 ggtaaaccatcattggttatttgtcttgattccggagcaggtaattatgagcagttatgg 600 Query: 541 Sbjct: 21620 ggtaaaccatcattggttatttgtcttgattccggagcaggtaattatgagcagttatgg 21679

mandiamantimiki kiliki minimia matuliiti

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59

Query: 661 Sbjct: 2174	ggcgttcattctgggagcgccagtggtatagtggcagacagtttcagagtagctcggcaa 720
Query: 721 Sbjct: 2180	ttgatcagcaggatagaggacgaaaacaccggagagataaaattacctcagttgtattgt 780
Query: 781 Sbjct: 2186	gatatteetgatgagagaataaaacaagegaaacaatgtgeggaaattetaggtgaacaa 840
Query: 841 Sbjct: 21920	gtttatagcgaatttccatggatagattctgccaaacccgttattcaagacaaacagcaa 900
Query: 901 Sbjct: 21980	ttaatattaaacagaacatggcgccctgccttgacggtgactggtgcagatgggtttcca 960
Query: 961 Sbjct: 22040	gcgatagctgatgcagggaacgtaatgcgccctgttacgtctttgaaattatccatgcgc 1020
Query: 1021 Sbjct: 22100	cttccaccactggttgatccagaagcagcttctgttgctatggaaaaagccctgacccaa 1080
Query: 1081 Sbjct: 22160	aaccctccctataatgcaaaggttgattttaaaatacaaaatggagggtccaagggatgg 1140
Query: 1141 Sbjct: 22220	aatgctcctttgctttccgattggttagcgaaagcggcatctgaagcatcaatgacttat 1200
Query: 1201 Sbjct: 22280	tatgataaacctgctgcttacatgggagaggggggcaccattccatttatgagtatgcta 1260
	ggcgagcaatttcccaaagcacaatttatgataactggtgttttaggcccccattccaat 1320
	gctcatggtccgaacgagttcttacatttggacatggtaaaaaaactcacctcatgtgtc 1380
	ccgtacgttctttatagtttttcacagaaaaataa 1416

TBLASTN 2.2.6 [Apr-09-2003]

Query= 1865.3 CONTIG=Contig42 POSCDS1=76674 POSCDS2=77765 SENS=p, Seq Id : 623

(367 letters)

Database: contigsLpPhiladelphia

51 sequences; 3,410,887 total letters

Searching.done

Sequences producing significant alignments:

Score E (bits) Value

LpPhiladelphia Contig49

718 0.0

>LpPhiladelphia Contig49 Length = 376826

Score = 718 bits (1853), Expect = 0.0Identities = 366/366 (100%), Positives = 366/366 (100%) Frame = +1

GNIMSPSIVFTGGGTAGHVTPNIALIKEFRKEGWNVEYIGSVSGIEKEMIEPLDIPFHGV Query: 1 GNIMSPSIVFTGGGTAGHVTPNIALIKEFRKEGWNVEYIGSVSGIEKEMIEPLDIPFHGV 60

Sbjct: 20005 GNIMSPSIVFTGGGTAGHVTPNIALIKEFRKEGWNVEYIGSVSGIEKEMIEPLDIPFHGV 20184

Query: 61 SSGKLRRYFSLKNLLDPFKIVLGIIQSSLLFYKIKPDVVFSKGGFVAFPVVVGAWLNRIP

SSGKLRRYFSLKNLLDPFKIVLGIIQSSLLFYKIKPDVVFSKGGFVAFPVVVGAWLNRIP 120 Sbjct: 20185 SSGKLRRYFSLKNLLDPFKIVLGIIQSSLLFYKIKPDVVFSKGGFVAFPVVVGAWLNRIP 20364

VVAHESDMSPGLANRLSFPFVNKICLTFDAGKKYFKRQDKIEVTGTPIRQQLLTGNRMKG Query: 121

VVAHESDMSPGLANRLSFPFVNKICLTFDAGKKYFKRQDKIEVTGTPIRQQLLTGNRMKG

Sbjct: 20365 VVAHESDMSPGLANRLSFPFVNKICLTFDAGKKYFKRQDKIEVTGTPIRQQLLTGNRMKG 20544

LELCGFNSSKPCLLVVGGSLGAGSINSCIRSALKQLTSEFQVIHLCGKGKLDSSLVGVEG Query: 181 LELCGFNSSKPCLLVVGGSLGAGSINSCIRSALKQLTSEFQVIHLCGKGKLDSSLVGVEG 240

Sbjct: 20545 LELCGFNSSKPCLLVVGGSLGAGSINSCIRSALKQLTSEFQVIHLCGKGKLDSSLVGVEG 20724

YCQFEYANEELADLFAASSVVISRAGANSLYEILALGKPHILIPISSQVSRGDQIQNARY Query: 241 YCQFEYANEELADLFAASSVVISRAGANSLYEILALGKPHILIPISSQVSRGDQIQNARY

Sbjct: 20725 YCQFEYANEELADLFAASSVVISRAGANSLYEILALGKPHILIPISSQVSRGDQIQNARY 20904

FQGLGISVVIQDELLKADVLLQAVQDVMRKKDEIDNKIKALKIESATDKIVAIIKEQAHV Query: 301

FQGLGISVVIQDELLKADVLLQAVQDVMRKKDEIDNKIKALKIESATDKIVAIIKEQAHV Sbjct: 20905 FQGLGISVVIQDELLKADVLLQAVQDVMRKKDEIDNKIKALKIESATDKIVAIIKEQAHV 21084

Query: 361 QTPRIV 366 QTPRIV

Sbjct: 21085 QTPRIV 21102

Database: contigsLpPhiladelphia Posted date: Nov 20, 2003 10:38 AM Number of letters in database: 3,410,887 Number of sequences in database: 51

Lambda

0.321 0.139 0.399

Gapped

Lambda K

> 0.267 0.0410 0.140

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61

Matrix: BLOSUM62 Query= 1865.3 CONTIG=Contig42 POSCDS1=76674 POSCDS2=77765 SENS=p (1092 letters) Database: /home/Gmp/rusniok/projets/legionella/pourBrevet-191103/contigsLpPhiladelphia 51 sequences; 3,410,887 total letters Searching.....done Sequences producing significant alignments: Score Е (bits) Value LpPhiladelphia Contig49 2165 0.0 >LpPhiladelphia_Contig49 Length = 376826Score = 2165 bits (1092), Expect = 0.0 Identities = 1092/1092 (100%) Strand = Plus / Plus ${\tt atgagcccaagtattgttttaccgggggaggaactgccggacatgtaacgcctaatatc}$ Query: 1 60 Sbjct: 20014 atgagcccaagtattgtttttaccgggggaggaactgccggacatgtaacgcctaatatc 20073 $\tt gctttgattaaggaatttcgaaaagaaggctggaatgtagaatatatcggctctgtttcc\\$ Query: 61 120 Sbjct: 20074 gctttgattaaggaatttcgaaaagaaggctggaatgtagaatatatcggctctgtttcc 20133 ggaattgaaaaggagatgattgagccgctggacattccttttcatggggtcagtagcggtQuery: 121 180 Sbjct: 20134 ggaattgaaaaggagatgattgagccgctggacattccttttcatggggtcagtagcggt 20193 aaattgcgcaggtattttagtttgaagaacttgcttgatcctttcaaaattgttctggga Query: 181 វិលពីកើតិកើតិប៉ុស្តែប៉ុស្តែប៉ុស្តែប៉ុស្តែប៉ុស្តែបាយប្រជាពលវិទ្ធាប៉ុស្តែប៉ុស្តែបាយប្រជាពលវិទ្ធិប 240 Sbjct: 20194 aaattgcgcaggtattttagtttgaagaacttgcttgatcctttcaaaattgttctggga 20253 Query: 241 attattcaatcttctttgctattttataaaatcaaacccgatgtggttttttcaaaaggt Sbjct: 20254 attattcaatcttctttgctattttataaaatcaaacccgatgtggttttttcaaaaggt 20313 Query: 301 ${\tt ggctttgtagcctttcctgtggttgtaggcgcctggttaaatcgaattcctgttgtcgct}$ 360 Sbjct: 20314 ggctttgtagcctttcctgtggttgtaggcgcctggttaaatcgaattcctgttgtcgct 20373 catgagtctgatatgagcccaggacttgcgaatcgcctatcctttcctttcgtcaataaa Query: 361 420 Sbjct: 20374 catgagtctgatatgagcccaggacttgcgaatcgcctatcctttcctttcgtcaataaa 20433 atatgtcttacttttgatgctggcaaaaaatactttaagcgtcaggataaaatagaagtg Query: 421

Sbjct: 20434 atatgtcttacttttgatgctggcaaaaaatactttaagcgtcaggataaaatagaagtg 20493

480

Query:	481	acgggtactccaattcgtcaacagctattaactggaaatcgaatgaaaggattggagtta	540
Sbjct:	20494	acgggtactccaattcgtcaacagctattaactggaaatcgaatgaaaggattggagtta	20553
Query:	541	tgcggatttaattcctccaaaccttgcctgcttgtagtgggaggaagcttaggggctggt	600
Sbjct:	20554	tgcggatttaattcctccaaaccttgcctgcttgtagtgggaggaagcttaggggctggt	20613
Query:	601	tcaattaacagttgtattcgaagcgcattgaaacaattgacatcagaatttcaagtcatt	660
Sbjct:	20614	tcaattaacagttgtattcgaagcgcattgaaacaattgacatcagaatttcaagtcatt	20673
Query:	661	catctttgtggcaagggaaaacttgattcttcattggttgg	720
		${\tt catctttgtggcaagggaaaacttgattcttcattggttgg$	
Query:		tttgaatacgccaatgaagagttggctgatctgttcgctgcttcttctgtggtgatttct	780
		${\tt tttgaatacgccaatgaagagttggctgatctgttcgctgcttcttctgtggtgatttct}$	20793
Query:		cgagcaggagctaattctttgtatgaaatattagcattaggaaaaccacatatcttaatt	840
		${\tt cgagcaggagctaattctttgtatgaaatattagcattaggaaaaccacatatcttaatt}$	
Query:		ccaatctcttcacaagtaagcagaggagatcaaattcagaatgcaaggtacttccaggga	900
		ccaatctcttcacaagtaagcagaggagatcaaattcagaatgcaaggtacttccaggga	
Query:		ttgggaataagcgttgtgattcaggacgagttattgaaagctgatgttctattacaggca	960
,		ttgggaataagcgttgtgattcaggacgagttattgaaagctgatgttctattacaggca	
Query:		gtacaggacgtaatgcgaaaaaagatgaaatagataataaaatcaaagcattaaaaatt	1020
		gtacaggacgtaatgcgaaaaaagatgaaatagataataaaatcaaagcattaaaaatt	
Query:		<pre>gagtctgccactgataagattgtggcaattatcaaggagcaagca</pre>	
		$\tt gagtctgccactgataagattgtggcaattatcaaggagcaagca$	21093
-		aggattgtatga 1092	
Sbjct:	21094	aggattgtatga 21105	
			-

TBLASTN 2.2.6 [Apr-09-2003]

Query= 2066.5 CONTIG=Contig46 POSCDS1=56766 POSCDS2=57173 SENS=p, Seq Id: 732 (150 letters)

Identities = 408/408 (100%)

Database: contigsLpPhiladelphia 51 sequences; 3,410,887 total letters Searching.done Score Sequences producing significant alignments: \mathbf{E} (bits) Value LpPhiladelphia Contig49 323 4e-90 >LpPhiladelphia_Contig49 Length = 376826Score = 323 bits (828), Expect = 4e-90Identities = 149/149 (100%), Positives = 149/149 (100%) Frame = +1 ${\tt IMYLRLLALSALCFVTSPIWSFTCIYTLVKDNCWTDYDVTVDVIEDSTSKTLLTLTAPKG}$ Query: 1 IMYLRLIALSALCFVTSPIWSFTCIYTLVKDNCWTDYDVTVDVIEDSTSKTLLTLTAPKG Sbjct: 89377 IMYLRLLALSALCFVTSPIWSFTCIYTLVKDNCWTDYDVTVDVIEDSTSKTLLTLTAPKG 89556 ${\tt KSWARGTFNCEAAEGLRYVAQFSPVFWQNDVGKTYPALRNWYLPAKVNPGDLAWTIPVCF}$ Query: 61 KSWARGTFNCEAAEGLRYVAQFSPVFWQNDVGKTYPALRNWYLPAKVNPGDLAWTIPVCF Sbjct: 89557 KSWARGTFNCEAAEGLRYVAQFSPVFWQNDVGKTYPALRNWYLPAKVNPGDLAWTIPVCF 89736 Query: 121 PADFAQVPFPPNVAGNCKCNFKNIPDPKL PADFAQVPFPPNVAGNCKCNFKNIPDPKL Sbjct: 89737 PADFAQVPFPPNVAGNCKCNFKNIPDPKL 89823 Database: contigsLpPhiladelphia Posted date: Nov 20, 2003 10:38 AM Number of letters in database: 3,410,887 Number of sequences in database: 51 Lambda K 0.323 0.138 0.473 Gapped Lambda K Н 0.267 0.0410 0.140 Matrix: BLOSUM62 Query= 2066.5 CONTIG=Contig46 POSCDS1=56766 POSCDS2=57173 SENS=p (408 letters) Database: /home/Gmp/rusniok/projets/legionella/pourBrevet-191103/contigsLpPhiladelphia 51 sequences; 3,410,887 total letters Searching......done Score E Sequences producing significant alignments: (bits) Value LpPhiladelphia_Contig49 809 0.0 >LpPhiladelphia_Contig49 Length = 376826Score = 809 bits (408), Expect = 0.0

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Strand = Plus / Plus

```
Query: 1
           \tt gtgactagcccaatttggtctttcacatgcatctatactttggttaaagacaattgttgg
            60
  Sbjct: 89419 gtgactagcccaatttggtctttcacatgcatctatactttggttaaagacaattgttgg 89478
           {\tt actgattatgatgttactgtcgatgtcattgaagattctacgtcaaaaactttgttgaca}
 Query: 61
            Sbjct: 89479 actgattatgatgttactgtcgatgtcattgaagattctacgtcaaaaactttgttgaca 89538
           \tt cttaccgctcccaaaggaaaatcatgggctagaggtactttcaattgtgaggctgctgaa
 Query: 121
           180
 Sbjct: 89539 cttaccgctcccaaaggaaaatcatgggctagaggtactttcaattgtgaggctgctaaa 89598
           gggttgagatatgtcgctcaattttcgcctgtcttttggcaaaatgatgttggaaaaact
 Query: 181
           240
 Sbjct: 89599 gggttgagatatgtcgctcaattttcgcctgtcttttggcaaaatgatgttggaaaaact 89658
           {\tt tacccggcattaagaaattggtatttaccagcaaaagtgaatcctggagatttggcctgg}
 Query: 241
           300
 Sbjct: 89659 tacccggcattaagaaattggtatttaccagcaaaagtgaatcctggagatttggcctgg 89718
           {\tt actatcccggtttgttttccggcagattttgctcaagttccctttccacctaatgtagca}
 Query: 301
           Sbjct: 89719 actatcccggtttgttttccggcagattttgctcaagttccctttccacctaatgtagca 89778
           ggaaactgtaagtgcaacttcaagaacattcctgatcccaagctttaa 408
 Query: 361
           Sbjct: 89779 ggaaactgtaagtgcaacttcaagaacattcctgatcccaagctttaa 89826
 .....
TBLASTN 2.2.6 [Apr-09-2003]
Query= 3159.2 CONTIG=Contig46 POSCDS1=34563 POSCDS2=35318 SENS=p, Seq
Id : 1433
       (265 letters)
Database: contigsLpPhiladelphia
         51 sequences; 3,410,887 total letters
Searching.done
Sequences producing significant alignments:
                                                   Score
                                                          Ē
                                                   (bits) Value
LpPhiladelphia_Contig49
                                                          537
                                                              e - 154
>LpPhiladelphia_Contig49
       Length = 376826
Score = 537 bits (1383), Expect = e-154
Identities = 264/264 (100%), Positives = 264/264 (100%)
Frame = +1
```

Query: 1	CVVESFFLILLFPMWKILYQLASPKNFYNYAGRLIPWLAVSALTTMAIGMV	
Sbjct: 671	CVVESFFLILLFPMWKILYQLASPKNFYNYAGRLIPWLAVSALTTMAIGMV 77 CVVESFFLILLFPMWKILYQLASPKNFYNYAGRLIPWLAVSALTTMAIGMV	WGLVFAPPD 60 WGLVFAPPD 67356
. Query: 61	YOOGDAYRIIFVHVPSAFISMALVAWACELATITIVA	
Sbjct: 673.	YQQGDAYRIIFVHVPSAFLSMALYAWMGFLAILLLVWRIKMAGLLIHKVAQI 57 YQQGDAYRIIFVHVPSAFLSMALYAWMGFLAILLLVWRIKMAGLLIHKVAQI	GACMAFLA 120 GACMAFLA GACMAFLA 67536
Query: 121	LITGSIWGKPMWGAWWVWDARLTSELTLLLYLAILAMWGAWW	
Sbjct: 675	LITGSIWGKPMWGAWWVWDARLTSELILLLLYLAILATYQAVKNKEDGDKII TITGSIWGKPMWGAWWVWDARLTSELILLLLYLAILATYQAVKNKEDGDKII	AILALVGL 180 AILALVGL AILALVGL 67716
Query: 181	IDLPIIHYSVYWWNTLHOGATLSVEAKDKIALSMIVDIX TOTA	
Sbjct: 6771	IDLPIIHYSVYWWNTLHQGATLSVFAKPKIALSMLYPLLITLLGFFLYSLWI 7 IDLPIIHYSVYWWNTLHQGATLSVFAKPKIALSMLYPLLITLLGFFLYSLWI	ILEKARNE 240 ILEKARNE ILEKARNE 67896
Query: 241	VLFRERKQSWVKIOFEEESDESVF 264	
Sbjct: 6789	VLFRERKQSWVKIQFEEESDESVF 7 VLFRERKQSWVKIQFEEESDESVF 67968	
Number of Number of Number of Number of Lambda 0.330 Gapped Lambda 0.267 (Matrix: BLOS Query= 3159. (75 Database: /h 191103/conti	С Н 0.0410 0.140	
	done	
		_
	oducing significant alignments:	Score E (bits) Value
LpPhiladelphi		1499 0.0
>LpPhiladelph Len	nia_Contig49 gth = 376826	
Score = 1499 Identities = Strand = Plu	bits (756), Expect = 0.0 756/756 (100%) s / Plus	
Query: 1	atgtggaagatattgtatcagttggcatcgccaaaaaatttttataactacgcgg	
Sbjct: 67216		ggacgt 60 ggacgt 67275
_	ctcattccctggttggcagtcagtgctttgactaccatggccattggtatggttt 	gggga 120
	tcattccctggttggcagtcagtgctttgactaccatggccattggtatggttt	gggga 67335

WO 2005/049642

Query: Sbjct:		ttggtatttgctccaccagattatcagcaaggggatgcataccgaattatttttgttcat	67395
Query:		gtacccagcgcttttttatcaatggcattgtatgcctggatggggtttctggccatttta	240
Sbjct:	67396		67455
Query:		ttgttggtgtggcgtatcaaaatggcagggcttttgattcataaggtcgcgcaattaggt	300
		ttgttggtgtggcgtatcaaaatggcagggcttttgattcataaggtcgcgcaattaggt .	
-	301 67516	gcctgcatggcatttcttgctttaattacagggagcatttggggtaaacccatgtggggt	360 67575
Query:	361	gcctggtgggtatgggatgcccgcctgacctcagaattaatacttttgttgctctatctg	420
Sbjct:	67576	gcctggtgggtatgggatgcccgcctgacctcagaattaatacttttgttgctctatctg	67635
Query: Sbjct:		gcaattctggctacctatcaagcggtaaaaaataaagaagatggagataaaataatagca	480 67695
Query:	481	attttagctttggtgggtttaattgatttaccaataattcattattcagtttattggtgg	540
Sbjct:	67696		67755
Query:	•	aatactttacaccaaggtgcaactttatctgtgtttgccaaacccaaaattgctctcagt	600
Query:	601	atgttgtatccattgttaatcactttqctqqqttttttcttqtattccttatqqatcatt	
		algetigtatecatigttaateattigtigggttttteetigtatteettatiggateatt	660 67875
Query:	661	ttggaaaaagcacgtaatgaagtcttattcagggagagaaagcaatcatgggttaagatt	720
Sbjct:	67876	ttggaaaaagcacgtaatgaagtcttattcagggagagaaagcaatcatgggttaagatt	67935
Query: Sbjct:		caatttgaggaagagtctgatgaatcagttttttga 756	
	. 	· ·	

TBLASTN 2.2.6 [Apr-09-2003]

Query= 4774.1 CONTIG=Contig46 POSCDS1=50654 POSCDS2=50950 SENS=m, Seq Id : 2523

(103 letters)

Database: contigsLpPhiladelphia

51 sequences; 3,410,887 total letters

Searching.done

Sequences producing significant alignments:

Score \mathbf{E} (bits) Value

LpPhiladelphia_Contig49

205 4e - 55

>LpPhiladelphia_Contig49 Length = 376826

Score = 205 bits (522), Expect = 4e-55Identities = 102/102 (100%), Positives = 102/102 (100%) Frame = -1

RRSKMPEIHTLDNPYITILTIFVLACFVGYYVVWKVTPALHTPLMSVTNAISSIIILGAL Query: 1 RRSKMPEIHTLDNPYITILTIFVLACFVGYYVVWKVTPALHTPLMSVTNAISSIIILGAL

Sbjct: 83615 RRSKMPEIHTLDNPYITILTIFVLACFVGYYVVWKVTPALHTPLMSVTNAISSIIILGAL 83436

IAAGSELIGCITWLGGIAIFITSINIFGGFVVTQRMLRMYKK IAAGSELIGCITWLGGIAIFITSINIFGGFVVTQRMLRMYKK Sbjct: 83435 IAAGSELIGCITWLGGIAIFITSINIFGGFVVTQRMLRMYKK 83310

Database: contigsLpPhiladelphia Posted date: Nov 20, 2003 10:38 AM Number of letters in database: 3,410,887 Number of sequences in database: 51

K Lambda 0.442 0.330 0.144

Gapped

Lambda

0.267 0.0410 0.140

Matrix: BLOSUM62

Query= 4774.1 CONTIG=Contig46 POSCDS1=50654 POSCDS2=50950 SENS=m (297 letters)

Database: /home/Gmp/rusniok/projets/legionella/pour Brevet-191103/contigsLpPhiladelphia 51 sequences; 3,410,887 total letters

Searching.....done

Score Sequences producing significant alignments: (bits) Value

LpPhiladelphia Contig49

589 e-169

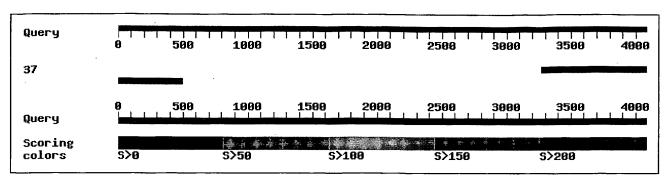
E

>LpPhiladelphia_Contig49 Length = 376826

Score = 589 bits (297), Expect = e-169Identities = 297/297 (100%) Strand = Plus / Minus

Query:	1	atgcctgaaattcatacacttgataatccttatattacaatattaaccattttcgtactg	60
Sbjct:	83603	atgcctgaaattcatacacttgataatccttatattacaatattaaccattttcgtactg	83544
Query:	61	<pre>gcctgttttgtaggttattatgtggtttggaaagtaacaccggctttacataca</pre>	120
Sbjct:	83543	gcctgttttgtaggttattatgtggtttggaaagtaacaccggctttacataca	83484
Query:	121	atgtcagtaaccaatgccatatccagtattattatacttggtgctttaattgctgcagga	180
Sbjct:	83483	atgtcagtaaccaatgccatatccagtattattatacttggtgctttaattgctgcagga	83424
Query:	181	agtgaattgatcggatgcataacctggttaggtggcatagccatattcattacttcaatt	240
Sbjct:	83423	agtgaattgatcggatgcataacctggttaggtggcatagccatattcattacttcaatt	83364
Query:	241	aatatttttggtggctttgtagtaactcaacgcatgcttcgcatgtataaaaaataa 297	•
Sbjct:	83363	aatatttttggtggctttgtagtaactcaacgcatgcttcgcatgtataaaaaataa 833	07

Exemple 5: Other examples of alignment fo sequences



>37 Length = 67519Score = 1439 bits (726), Expect = 0.0Identities = 784/802 (97%), Gaps = 1/802 (0%) Strand = Plus / Plus Query: 3275 3334 Query: 3335 ttggacttgctatgtgtcattccactgggtcaattggcgacacactgattggccctttct Sbjct: 63980 ttggacttgctatgtgtcattccactgggtcaattggcgacacactgattggccctttct 64039 Query: 3395 $\verb|attatcctgaaatcctgacaagagctctctatggcttaatctataagctgcttgtgatta|\\$ Sbjct: 64040 attatectaaaaccetgacatgaactetetatggettaatetataagetgettgtgatta 64099 atttcatcgcaatataagccattaaaataccgctaagtaactctattttttgccatactt 3514 Query: 3455 Sbjct: 64100 atttcatcgcaatataagccattaaaataccactaagtaactctatttttttgccatactt 64159 Query: 3515 tattttgagttaacaggtttgaaaaataacgagtagtcatcgttaatgaactgaaccaaa 3574 Sbjct: 64160 tattttgggttaataggtttgaaaaataacgagtagtcatcgttaatgaactgaaccaaa 64219 tcatgctggcagaaattacccctgctaaaaaagccagtttatggtcaggatattgactgc 3634 Query: 3575 Sbjct: 64220 tcatgctggcagaaattacccctgctaaaaaagccagtttatggtcaggatattgactgc 64279 Query: 3635 tgccgctacccacaatcaccagggtgtctataatggcgtgaggattaagcagactaaacc 3694 Sbict: 64280 tgccgctgcccacaatcaccagggtatctataatggcgtgaggattaagcagactaaacc 64339 Query: 3695 ccagggcaaataaaatgatttgcattcttgtatgaqgctgatgtgtttctacaacggttt Sbjct: 64340 ccagggcaaataaaatgatttgcattcttgtatgaggctgatgttttctacaacggttt 64399 Query: 3755 gtttgtttttggacagcgcgctttttaagttttttattgcataataaattaaaaggcag

Sbjct: 64400 gtttgtttttggacaacgcgctttttaagttttttattgcataataaattaaaaggcag 64459

PCT/IB2004/003578

Query: 3815 agcctagccataccatccagatttgcaagtttggatgagccaaaagcaattgatgtaaac 3874 Sbjct: 64460 agcctagccataccatccagatttgcaagtttggatgagccaaaagcaattgatgtaatc 64519 Query: 3875 3934 3994 Sbjct: 64580 ccgcatgattttttcgcgcaccttgcctgataagaaaaacattttgcggacctaaggcca 64639 Query: 3995 ttatcaaagataatcccaagagtaatccattaaaattaaatcaacataattgcatcaggat Query: 4055 agtaaaaaaaaaggcgattata 4076 111 1111111111111111 Sbjct: 64700 agt-aaaaaaaaggcgattata 64720 Score = 957 bits (483), Expect = 0.0Identities = 495/499 (99%)Strand = Plus / Plus Query: 1 $\verb|ctacaaattttgcaaggttattaaatagtggttttcatctggcggcctattgttttttg|\\$ 60 Sbjct: 63429 ctacaaattttgcaaggtaattaaatagtggttttcatctggcggcctattgttttttg 63488 Query: 61 ggaaagccataagcattctgccaattgatccatgattaaatgttcaacagccatqqqatc 120 Sbjct: 63489 ggaaagccataagcattctgccaattgatccatgattaaatgttcaacagccatgggatc 63548 Query: 121 ctggtatttttctattaacttggtgtaaacagtacggatgccttgtggcctatcggtcgt 180 Sbjct: 63549 ctgatatttttctattaacttggtgtaaacagtacggatgccttgtggcctatcggtcgc 63608 Query: 181 240 Sbjct: 63609 tacttgttcgcgaacggcaagatgaagccccatatgaaggaatggtttcgcctaa 63668 $\verb|ttcaggata| at a a gtat gttcaggaa a a gat t ga a t t t gttca a t gat t t a t gg t a t t c$ Query: 241 300 Sbjct: 63669 ttcaggataataagtatgttcaggaaaagatggaatttgttcaatgactttatggtattc 63728 · Query: 301 $\verb|cggatgatcaagaatcacttgggcaatttcttttccaagggagaaagttctttttatt|$ 360 Sbjct: 63729 cggatgatcaagaatcacttgggcaatttctttttccaagggagaaagttctttttatt 63788 Query: 361 ctggtacttattccagctgataaaaaatagctgtcgagtttcttgtactgtatcgccgta 420 Sbjct: 63789 ctggtacttattccagctgataaaaaatagctgtcgagtttcttgtactgtatcgccgta 63848

```
Query: 421
            {\tt aaacataatggcccgattgatataaaatgatccattttaactgaataaaaaagtaacaac}
             4144144141111414141414141414141414444
Sbjct: 63849 aaacataatggcccgattgatataaaatgatccattttaactgaataaaaagtaacaac 63908
Query: 481
            aatgttgatgtgcaaatat
             111111111111111111111
Sbjct: 63909 aatgttgatgtgcaaatat 63927
  Database: /local/http/htdocs/IPF_Gmp/legionella/blastdb/contig
   Posted date: Sep 10, 2003 12:44 PM
  Number of letters in database: 3,410,887
 Number of sequences in database: 51
Lambda
          K
   1.37
           0.711
                     1.31
Gapped
Lambda
          K
   1.37
           0.711
                     1.31
Matrix: blastn matrix:1 -3
```

Alignment of a portion of contig48 (Seq Id 48) of the Paris strain with all the contigs of the Philadelphia strain.

The positions of this fragment in the contig are indicated in the line starting with « Query= ». In this example, the first position (noted 1 in the alignment) of the fragment is thus position 50760 in contig48 of the Paris strain. This fragment terminates in position 56335 in contig48. 50760 should thus be added to the position indicated by the alignment to have the position in the total sequence of the contig. The positions in the contig of the Philadelphia strain are unchanged.

In this exemple, nous pouvons voir que the regions of the Paris strain between the positions 561(+50760) and 2096(+50760) and between 2622(+50760) and 3981(+50760) are absent from the Philadelphia strain. These regions thus contain the following ORFs, specific to the Paris strain:

```
322.3 (Seq ID 1466)
321.3 (Seq ID 1462)
3005.2 (Seq ID 1329)
5208.1 (Seq ID 2782)

BLASTN 2.2.6 [Apr-09-2003]

Reference: Altschul, Stephen F., Thomas L. Madden, Alejandro A. Schaffer, Jinghui Zhang, Zheng Zhang, Webb Miller, and David J. Lipman (1997), "Gapped BLAST and PSI-BLAST: a new generation of protein database search
```

72

programs", Nucleic Acids Res. 25:3389-3402.

Query= Lp Paris Contig48_50760-56335 (5576 letters)

Database: /local/http/htdocs/IPF_Gmp/legionella/blastdb/contig

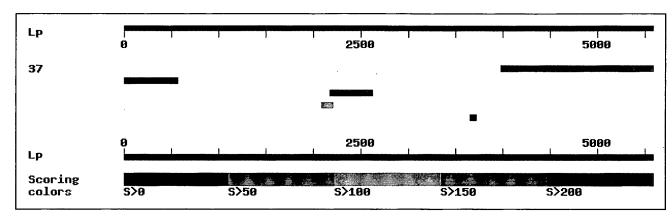
51 sequences; 3,410,887 total letters

Searching.done

Sequences producing significant alignments:

Score E (bits) Value

37 2892 0.0



>37

Length = 67519

Score = 2892 bits (1459), Expect = 0.0 Identities = 1561/1595 (97%) Strand = Plus / Plus

Query: 3982 cttatggcaaatatttatccctcaaagcgtttctcaataagccattgttaccatgaacca 4041

Sbjct: 50163 cttatggcaaatatttatccctcaaaatgtttctcaataagtcattgttaccttgaacca 50222

Query: 4042 gggtaagctttaattttcttaaacaaaatggagtatttagtcctcccttagatggaagct 4101

Sbjct: 50223 gggtaagctttaatttcttaaacaaaatggagtatatagtcctctcttagatggaagct 50282

Query: 4102 ctttcactgctttgctgagcaaatatttgacgctcctcactgattaattcaatccatttt 4161

Sbjct: 50283 ctttcactgctttgctgagcagatatttgacgctcctcactgattaattcaatccatttt 50342

Query: 4162 ttaggtgtcatgtcgttttgtatcaataagttgatcgctgcactaggatattgactttgt 4221

Query: 4222 gtaatgacaaaataaaagtaagcggtcaatttgcttctgcataaagcccaaccacttttt 4281

Sbjct: 50403 gtaatggcaaaataaaagtaagcggtcaatttgcttctgcataaagcccaaccacttttt 50462

Query: 4282 gcgacacattgttgcaaactagcaaattcaatttgatcttctttgactaattgttgcagc 4341

Sbjct:	50463	gcgacacattgttgcaaactaggaaattcaatttgatcttcttttgactaattgttgcagc .	50522
Query: Sbjct:		ttttgaatcgtgtcagggatacatgcatcaaaaaacaatgtggtcccttgctgtccccat	4401 50582
Query: Sbjct:		acgtcacttaatataattctttttaaaatgttcatgtcacaaatttgttgctcatgggga	4461 50642
Query: Sbjct:		ataggtttctcaattctatgtgttaccggttgccccaacatggaatgtgtccattgctga	4521 50702
Query: Sbjct:		taataagcccgaccgtaaattaatataccacccaaaataaaatcaaaggcagttggggct	4581 50762
Query: Sbjct:		gcataccaaagagcagtgaagaatttggatttgatgacgtccactttggggtagggatac	4641 50822
Query: Sbjct:		tgatggttattggttttaatgatatgtttgcgatcgtattgcatttttaaaggttcatat	4701 50882
Query: Sbjct:		ttaaccaaggatttagcctggtattgattctggccaatacttatcgtttcttctttt	4761 50942
Query: Sbjct:		ggtattcctccatcgactattgctaaatgaaagttatgatttaaaacatgttcataaccc	4821 51002
Query: Sbjct:		agaaaatgacaatcaggatttttagataaggtatttgctaaatgttgtcgcaatgcctcc	4881 51062
Query: Sbjct:		ccgactttaatattaggatcataaaaatctggatgtttggctaaattcattacccaaaat	4941 51122
Query: Sbjct:		acattgactttattcttttcaccagcaggaatatctcgtaacatcattccccaacaatac	5001 51182
Query: Sbjct:		ccaataacttcgtcgtttttatccttggcaataaaacaaatagtctctttataatgtaac	5061 51242
Query:	5062	attgaagataggacgatatcccctataaaaggcaaatcgccaaaatttgctccatgtgct	5121

Sbjct: 5	51243	attgaagataggacgatatcccctataaaaggcaaatcgccaaaatttgctccatgggct	51302
Query: 5		aaagaatcgatacgtttaatttcacccattgctttattaagttcagcagtattattggga 	5181 51362
Query: 5		tctaaaatctgaacttctttgaagccatagggcttgccttccatcatttttaacgaaaaa	5241 51422
Query: 5		ttaaaggtacattcgccatctaactttttgtggtgtttaattttaagaggattttgattt	5301 51482
Query: 5		ctgcaatttaacagatctgtccataattgatgaatgtaaagtaatctttcatcatttggt !	5361 51542
Query: 5		gaggtatttcgcaaattgattaattgctgggtgaaacgatgcacatattctgtatggaga !!	5421 51602
Query: 5		taaatagcatggctactgaaatcaacaacaaactctgccgtcattaattttttgatttcg 	5481 51662
Query: 5		ggtaagtgatattcaacaggatctgtatatgattttactattgcttctagtgaactccat	5541 51722
Query: 5		ttgcgttcaagtacatcgatttttgaatacgacat 5576 	
	cies =	bits (469), Expect = 0.0 543/569 (95%), Gaps = 9/569 (1%) s / Plus	
Query: 1 Sbjct: 4		atggcgaaatcattgtcgcaattagattctgctaatttgctccctgttttaacatggaa 	60 46599
Query: 6		caagcagaacgcattggaaaacaaatcaataagctcttacagcatgagttttgcgaggaa 	120 46659
Query: 1		aacatcaatccaaagaaatttgcctctatcagtcacaatatcctgcccaaaattatgaca 	180 46719
Query: 1		gaaacatttttaggagtaaccccgccagaaaactggcagcaattaagcgacgatattata	240

Sbjct: 46720 gaaacatttttaggagtaacccctccagaaaactggcagcaattaagcgacgatattata 46779
Query: 241 aaaaactgcatcgcaaacaagaatctatgcaaaaaagcagctcgcaaagagctggaagaa 300
Query: 301 tgcatcaaaccgagaattcctttgatcctgatacaatttggcccctggcttgctcaaaat 360
Query: 361 tgtccacaattgaataagtctttgattgaacaatggccaaataaacaggctactctcaaa 420
Query: 421 aagataattaatgaaaacaaaagtgccgagtaatcgagacaggcttaattacgagagtta 480
Query: 481 tgcctgataaaaccacatttatttacctaatttcatcaaatataactcacc 531
Query: 532 gatatgatctacaaccaagttcttaaaac 560
Score = 656 bits (331), Expect = 0.0 Identities = 423/451 (93%), Gaps = 2/451 (0%) Strand = Plus / Plus
Query: 2172 tatatggccgataaaatttgccagggtcaataaatagtattctgatggttaggtaataaa 2231
Query: 2232 tgatgaagagttcttttggtgaaatgaataaaagaccccttttttattgagcgactctta 2291
Query: 2292 aaaa-gccatttgctttattctgtgcttttggcaagtgacatgatcgcattcaggttttg 2350
Query: 2351 tttacatgctaggtgtggttttttcccgaggcggttgtgagtttgataccataggtttta 2410
Query: 2411 ttgaataggcgccgcttgggaaaaaacaattgtcagtaaggattgcccctgtaatcagat 2470
Query: 2471 gggtaagccgattggcttcggcacatctcgaatcagtgacaacctctttcgctttttcat 2530

```
Sbjct: 48226 gggtaagccgattggcttcggcacatctcgaatcagtggcaacctctttcgctttttcat 48285
Query: 2531 ctttatttcgcataacaatcctgtgaagttaatctttgcagaggacaccatgatggtttc 2590
           Sbjct: 48286 ctttatttcgcataacaatcctgtgaagttaatctttgcagaggacactatgatggtttc 48345
Query: 2591 atgtcatacaaacgaagcaaccggataccga 2621
           +11111111111111111111111111111111111
Sbjct: 48346 atgtcatacaaacgaagcaaccgggtaccga 48376
 Score = 147 bits (74), Expect = 9e-35
 Identities = 98/106 (92%)
 Strand = Plus / Plus
Query: 2097 acctttctggagtttcccaactacaagatgatactgcgttataataactccatttattat 2156
           Sbjct: 44922 accttcctggcgtttcccaactacaagatgatcctacgttataataactccatttattat 44981
Query: 2157 actggggctatcgagtatatggccgataaaatttgccagggtcaat 2202
            Sbjct: 44982 gccggggctatcgggtatatggctgataaaatttgccagggtcaat 45027
 Score = 109 bits (55), Expect = 2e-23
 Identities = 64/67 (95%)
 Strand = Plus / Plus
Query: 3653 cgaatccttacaggaaaacgaagcttatggaagtccaacaaggaagaggtagtaaattca 3712
           Sbjct: 48378 cgaatccttacaggaaaacgaagcttatggaaggccaacaaggaacagggagtaaattca 48437
Query: 3713 taacgcc 3719
           111111
Sbjct: 48438 taacgcc 48444
 Database: /local/http/htdocs/IPF Gmp/legionella/blastdb/contig
   Posted date: Sep 10, 2003 12:44 PM
 Number of letters in database: 3,410,887
 Number of sequences in database: 51
Lambda
         0.711
   1.37
                  1.31
Gapped
Lambda
   1.37
          0.711
                  1.31
Matrix: blastn matrix:1 -3
```

Alignment of a portion of the contig45 (Seq Id 45) of the Paris strain with all the contigs of the Philadelphia strain.

The positions of this fragment in the contig are indicated in the line beginning with « Query= ». In this example, the first position (noted 1 in the alignment) of the fragment is thus position 44722 in the contig45 of the Paris strain. This fragment terminates at position 52680 in the contig45. It is thus required to add 44722 à to the indicated by the alignment to have the position in the total sequence of the contig. The positions in the contig of the Philadelphia strain unchanged.

In this example, we can see that the region of the Paris strain between the positions 1333(+44722) and 6899(+44722) is absent from the Philadelphia strain. This region thus contains the following ORFs, specific to the Paris strain:

```
4927.1 (Seq ID 2623)
413.5 (Seq ID 2069)
415.2 (Seq ID 2087)
417.3 (Seq ID 2102)
```

BLASTN 2.2.6 [Apr-09-2003]

Reference: Altschul, Stephen F., Thomas L. Madden, Alejandro A. Schaffer, Jinghui Zhang, Zheng Zhang, Webb Miller, and David J. Lipman (1997), "Gapped BLAST and PSI-BLAST: a new generation of protein database search programs", Nucleic Acids Res. 25:3389-3402.

Query= Lp Paris Contig45_44722-52680 (7959 letters)

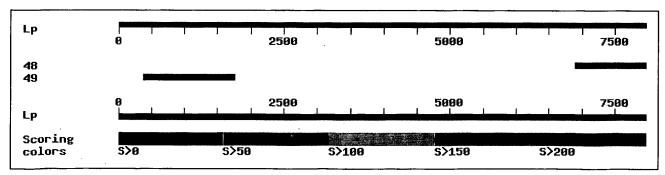
Database: /local/http/htdocs/IPF_Gmp/legionella/blastdb/contig 51 sequences; 3,410,887 total letters

Searching.done

Sequences producing significant alignments:

Score E (bits) Value

48 49 1943 0.0 94 2e-18



>48

Score = 1943 bits (980), Expect = 0.0
Identities = 1040/1060 (98%)
Strand = Plus / Minus

	·	
Query: Sbjct:	attctaatggtggccagaggcagaatcgaactgccgacacgaggattttcagtcctctgc	6959 55039
Query: Sbjct:	tctaccgactgagctatctggccactcaaggcttgctattaaactccgagatcgactttg	7019 54979
Query: Sbjct:	agtcaagtgttgattaagttttttttgaaaataatttttttt	7079 54919
Query: Sbjct:	ctgctcctagaaaagcgttgttttgataagctcaatattagcttccaggctcttgcctg	7139 54859
Query: Sbjct:	cagttaattcagcaatacgttttaacaggaagggggttaatgattttccgctcatgtgtt 	7199 54799
Query: Sbjct:	tggcttcttcatgagcctgttttatgtatggactgatttcctcatcagatagttccgctg	7259 54739
Query: Sbjct:	atactggaatggggtttgcgacgactattccgtttttcatgttcaatttctgttgaattg	7319 54679
Query: Sbjct:	acatgagatttgctacttcctcgaccgaatttaagcgttgtgggactggtattccactcg	7379 54619
Query: Sbjct:	atctgctgtaaaaagcagggaattcgtctgtggcataacctatgaccggcaccccaaacg	7439 54559
Query: Sbjct:	tttcaagaacttccaatgtttttggtaagtcgagaatcgattttgcgccagaacagacta 	7499 54499
Query: Sbjct:	cggtaactggcgtattggatagttctataagatcagctgaaatatcaaaactcattgtca	7559 5 4 439
Query: Sbjct:	cgtcttgatgaacaccacctatacctccggtgacaaatagggggagcctagccatatggg	7619 54379

```
Query: 7620 cacagaacatggttgctgctaccgttgtgctggcagtcactttacgagataatacaaaag 7679
            Sbjct: 54378 cacaaaacatggttgctgctaccgtcgtgctggcagtcactttacgagataatacaaaag 54319
 Query: 7680 aaatgtctctgcgagaggcttttattacttctttttgcagcgcgagatgctccataactt 7739
            Sbjct: 54318 aaatgtctctgcgagaggcttttattacttctttttgaagcgcgagatgttccataactt 54259
 Query: 7740 cttgagttaaaccaatacggattttcccttggtgcatcgctatagtagctggaatggcgc 7799
            រត់ពីពីពីពេយ៌យក់ ពីពីពីតែពេយ៍យើយពីពេលពីពេលពេយ៍យើយពីការីការីពេយ៍
 Sbjct: 54258 cttgagttaaaccaacaggattttcccttggtgcatcgctatagtagctggaatggcgc 54199
 Query: 7800 cttgtctacgaataatattttcaacttctattgccgtagttaaattatcagggtagggca 7859
            Sbjct: 54198 cttgtctacgaataatattttcaacctctattgccgtagttaaattatcagggtagggca 54139
           ttccatgagagataatggtcgactcaagagcaacaattgggtttttatcattgatagcat 7919
 Query: 7860
            រីសំពីស៊ីស៊ីស៊ីស៊ីស៊ីស៊ីសាលស៊ីស៊ីសមសាស៊ីស៊ីសាលស្រាស៊ីសាល
 Sbjct: 54138 ttccatgagagataatggtcgactcaagagcaacaattgggtttttatcattgatagcat 54079
 Query: 7920 ccagtacttcttcgttaaattccaacaagtcatgaaacat 7959
            Sbjct: 54078 ccagtacttcttcgttaaattccaacaagtcatgaaacat 54039
 >49
         Length = 376826
 Score = 93.7 bits (47), Expect = 2e-18
 Identities = 102/119 (85%), Gaps = 1/119 (0%)
 Strand = Plus / Plus
            {\tt atttgccctgtgtattgtttagtgttggtcgagcggttcactctctgttgaaacccggtanger} \\
Query: 1215
            អំណើរអំណើយ អំណើរណី ហើរដែលអាយ ដែរវិយាជាវិ
                                                                1274
aaa-ccgtaaagctcgaagaagggggcaaatcaatcgttataaggcaaacgatcccgcc
Query: 1275
            1332
Sbjct: 207743 aaagccataaagctcgaagtagggggcaaatcaatcgttataaggcaaacgatctcgcc 207801
  Database: /local/http/htdocs/IPF_Gmp/legionella/blastdb/contig
   Posted date: Sep 10, 2003 12:44 PM
  Number of letters in database: 3,410,887
  Number of sequences in database: 51
Lambda
         K
   1.37
         0.711
                  1.31
Gapped
Lambda
         K
   1.37
          0.711
                  1.31
Matrix: blastn matrix:1 -3
```

Alignment of a portion of contig39 (Seq Id 39) of the Paris strain with all the contigs of the Philadelphia strain.

The positions of this fragment in the contig are indicated in the line starting with « Query= ». In this example, the first position (noted 1 in the alignment) of the fragment is thus position 3990 in contig39 of the Paris strain. This fragment terminates at position 8972 in contig39. 3990 should this be added to the position indicated by the alignment to have the position in the total sequence of the contig. The positions in the contig of the Philadelphia strain are unchanged.

In this example, we can see that the region of the Paris strain between the positions 1264(+3990) and 4465(+3990) is absent from the Philadelphia strain. This region thus contains the following ORFs, specific to the Paris strain:

```
3396.1 (Seq ID 1588)
3395.2 (Seq ID 1587)
3394.1 (Seq ID 1586)
```

BLASTN 2.2.6 [Apr-09-2003]

Reference: Altschul, Stephen F., Thomas L. Madden, Alejandro A. Schaffer, Jinghui Zhang, Zheng Zhang, Webb Miller, and David J. Lipman (1997), "Gapped BLAST and PSI-BLAST: a new generation of protein database search programs", Nucleic Acids Res. 25:3389-3402.

Query= Lp Paris Contig39_3990-8972 (4983 letters)

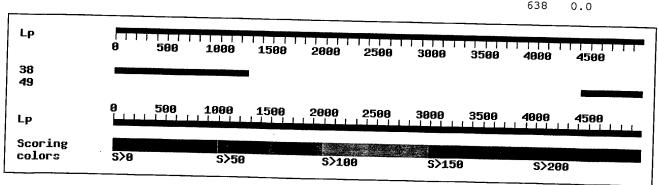
Database: /local/http/htdocs/IPF_Gmp/legionella/blastdb/contig 51 sequences; 3,410,887 total letters

Searching.done

Sequences producing significant alignments:

Score E (bits) Value

2123 0.0 638 0.0



>38

38

Score = 2123 bits (1071), Expect = 0.0 Identities = 1215/1263 (96%) Strand = Plus / Minus

Query: 1 Sbjct: 12277	ctttggttctacatgagcttgcctgagatgtttgcccttatcgattgcaacaatttttac 60
Query: 61 Sbjct: 12217	gccagttgtgagcgtttgtttcgtcctgatttaaaggacgtccccatcgtggtgctatcc 120
Query: 121 Sbjct: 12157	aataatgatggctgttgtatcgcacgctcgaatgaagccaaagcattgggcattgccatg 180
Query: 181 (ggcgagccgtacttcaaaattaaacatttgtgcaaacagcatggagtgaaagctttttcc 240
Query: 241 t Sbjct: 12037 t	caaattatacgctgtatggcaacatgagtcatcgtgtgatgtgcactattgaagaagcc 300
	ggccccatatagaagtttactcgattgatgaagcgtttcttgatttaaggagtttaccg 360
	ttgatagccatgattcgttttgcgagcagttacaaaagaaaatcttgaagcacacagga 420
Query: 421 a Sbjct: 11857 a	tacccacttccatcggtattggacctactaaaacactagctaaagccgccaatcattta 480
Query: 481 to	gcaaaaaagtttataaaatccctgtgtttaatatcacctcgaatcgtgggcggttattg 540
	acagatttccgttggggacatttggggagtagggcggcaatgggccaataaattaatt
Query: 601 tc	gcgaggcattcatacggcttatgatttggcaatgaccaatcctcaccttctgaagaaa 660
Query: 661 tg	ttttaacgtcgtgttgatgcgtaccgccatggagcttcaaggaattgcttgtggcggt 720

Query: Sbjct:		ttagaggcaatagagcctaagcaaagtattatgtcatctaaaagttttggtcagatgcaa 	780 11498	
		J JJ J J J J J J J J J J J J J J J J J		
Query:		actcaacttgcttcgattgaggaatcaatcagtagccattgtgcccgtgcggtggagaaa 	840	
Sbjct:	11497	actcaaattgcttcgattgaggaatcaatcagtagccattgtgctcgtgcggtggagaaa 1	11438	
Query:	841	atgcgtcgccagcaattggtggcgaagcgcctggttgtatttgtgcatacgaaccgattt	900	
Sbjct:	11437	atgcgtcgccaacaattggtggcgacgcgtctggttgtatttgtgcatacgaaccgattt 1	11378	
Query:	901	cgcgaagatttggcacagcactttcagtccatcgaatttaagctgattaatcctacagat	960	
Sbjct:	11377		11318	
Query:	961	gatttgcgcttaattaccaaaatggccaagcgatgtctgcaacgcatttttaaaccaggg	1020	
Sbjct:	11317	gatttgcgcttaattaccaaaatagccaaaagatgtctgcaacgcatttttaaaccaggg 1	11258	
Query:	1021	tattactataaaaaggcaggagtatgtcttgaggacttaattcccaaaaacccacgacag	1080	
Sbjct:	11257	tattactataaaaaggcaggggtatgtcttgaagacttaattcctaaaaaaccacgacag 1	11198	
Query:	1081	ctggatatgttttatcaaccaagtgacgagcatctaaaccacacggaacaattgatggcg	1140	
Sbjct:	11197	ctggatatgtttcatcaaccaagtgatgagcatctaaaacacaccgaacaattgatgggt 1	1138	
Query:	1141	gtctttgaccaaatcaatcaaaaatacggacgaagtacaatccgcctcgcggcagagggt	1200	
Sbjct:	11137	gtctttgaccaaatcaatcaaaagtatggaagaagcacgattcggttagccgccgaaggc 1	.1078	
Query:	1201	tattcaaaaccttgggcgatgcgtgctgaactgaaatcgcctgcct	1260	
Sbjct:	11077	tattcaaaaccctgggagatgcgtgctgagctgaaatcacctgcttataccacgcgatgg 1	1018	
Query:		tct 1263		
Sbjct:		tet 11015		
>49	Len	ngth = 376826		
Score = 638 bits (322), Expect = 0.0 Identities = 469/518 (90%) Strand = Plus / Plus				
Query:	4466	gaacaataatcactgataaaaagatcttgagcaaaagtctcaaaatcaaaatagcagatc	4525	
Sbjct:	203775		203834	
Query:	4526	aaactatccggcatttgatgggcataacagtcatcaaataattgcctggcaaaatcaacc	4585	

Sbjct: 203835 aaactatccggtatttgatgggcataacagtcatcaaataactgtctggcaaaatcaacc 203894 tctgaatcataacaaccttggtaatgatcctccaacatggtttgtgcatcatctaccgag 4645 Query: 4586 Sbjct: 203895 tctgaatcatagcaaccttggtaatgatcctccaacatggtttgtgcatcttctaccgaa 203954 Query: 4646 Sbjct: 203955 taatcgcagagaagggctaagcctagctctccatgttcttgaataaatgaagcatactcc 204014 Query: 4706 acaatgttgcttattccttcatattcatgaattctcatgctgccgaaaccttcataatcg 4765 Sbjct: 204015 acaatattacttattgactcgtactcatggatttttgatgctgccgaatccttcataatcg 204074 Query: 4766 tggatggcaaattcctcagcattggtttctgggctattatccaacatttcccagatttct 4825 Sbjct: 204075 tgtatggcatattcttcagcattgggctcagggctgttatccagcatttcccagatttct 204134 Query: 4826 ttcatqatqtcatcttcqctttqaqtaqcatctatccatacaccatqcaqtatqqcqttq 4885 Sbjct: 204135 tccataatgtcatcttcgctttgtgtggcatctatccaaacaccatgcaggatggcattg 204194 Query: 4886 ttgtaagaggctaaacaggcgacgtagattgaaggggtqtccatgggattatctccttgt 4945 Sbjct: 204195 ttgtatgaggctaaacacgcgacgtagattgaaggggtgtccatgggattatttccttgt 204254 Query: 4946 attaagggagctatcccacacgggagcttgctcccgtg 4983 Sbjct: 204255 attaagggagcaatcccacacgggagcttactcccgtg 204292 Database: /local/http/htdocs/IPF Gmp/legionella/blastdb/contig Posted date: Sep 10, 2003 12:44 PM Number of letters in database: 3,410,887 Number of sequences in database: 51 Lambda K 0.711 1.37 1.31 Gapped Lambda 0.711 1.31

Matrix: blastn matrix:1 -3

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Example 6: RESULTS

See hereinbelow Tables V, VI, I and II hereinbelow.

Example 7: Sequencing and annotation of the genome of L. pneumophila Lens strain

Comparison of the sequences of the genomes of *L. pneumophila* Paris strain, Lens strain and Philadelphia strain (http://genome3.cpmc.columbia.edu/~legion/index.html), three strains of serogroup 1, shows that around 88% of these genomes are very strongly preserved (95 to 100% of proteic identity), whereas the remaining 12% are specific to each strain. These results suggest that there is a large genomic diversity at the very centre of the *L. pneumophila* species.

The Table XVI hereinbelow comprises for each of the ORFs identified in the Lens strain its position on the genome, the existence of a peptide signal, the best result of the blast on nrprot (Best-Blastp). The ORFs specific to the strain *L. pneumophila* Lens relative to the strain *L. pneumophila* Paris were identified in considering as specific the ORFs having a percentage of proteic homology less than 75%. In the case where the ORF is preserved in the two genomes, the percentage of homology between the two proteins is mentioned. Finally, the ORFs specific to the *Legionella* genre were identified in considering as specific the ORFs having a percentage of proteic homology with sequences of the bank nrprot less than 25 %.

In conclusion, these results help define DNA probes for developing a typing tool. The utilization of this tool on a large number of strains isolated from patients and strains isolated from the environment can confirm if this tool can predict the risk associated with a strain by definitely discriminating the strains isolated from patients of the other strains.

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Material supplied:

- The complete sequence of the genome of *Legionella pneumophila* Lens strain made up of the long chromosome of around 3.33 Mb and of a long plasmid of around 60 kb.
- A list of specific coding phases of *L. pneumophila* Lens strain annotated with their nucleotidic sequences.

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Materials and methods.

1. Construction of the shotgun bank of small fragments (size 1.5 to 2.5 kb)

The chromosomal DNA of the strains studied was prepared by a classic method including proteinase K treatment and phenol extraction (9). Around 40 ug of DNA were broken by nebulization (1 minute under pressure of 1 bar) (4). The ends of the fragments of DNA were rendered free by having the DNA-polymerase of the bacteriophage T4 act for 15 minutes at 37°C in the presence of the 4 tri-phosphate nucleotides. The enzyme was inactivated by incubation of 15 mn at 75°C. Adaptors (invitrogen Cat. N° 408-18) have then been ligatured to these ends. After ligature, the fragments of chromosomal DNA having a size between 1500 and 2500 base pairs were purified after electrophoresis on agarose gel. The vector utilized for construction of the bank, pcDNA2.1 (Invitrogen), was digested by the enzyme BstX1 and purified by geneclean (BIO-101) after electrophoresis on agarose gel. The chromosomal DNA and the purified vector were ligatured by action of the ligase of the bacteriophage T4. The ligation mixture was introduced by transformation in the strain of Escherichia coli XL2-blue (Stratagene). About 4000 colonies are obtained per ul of the ligation mixture.

2. Preparation of plasmids and sequencing

The plasmids were prepared from bacterial colonies with the «TempliPhi DNA sequencing template amplification» kit marketed by Amersham Bioscience. The chromosomal inserts were sequenced from their two ends by utilizing the universal primer T7 by following the recommendations of the supplier (Applied-Biosystems). The sequences were determined by utilizing automatic sequencers of type 3700 (Applied-Biosystem).

3. Assembling of sequences

The sequences were assembled by utilizing the set of software developed at the University of Washington, Phred, Phrap and Consed (5, 8). The sequence was finished by utilizing the set of CAAT-box software (7). The finishing stage corresponds to resequencing the regions where the sequence is only slightly secure and sequencing of the regions situated between the contigs. It was done either by sequencing PCR products or by operating on the clones of the bank. The oligonucleotidic sequences were defined by utilizing consed and Primo software (8, 10).

3. Annotation of sequences

The identification of the coding phases (CDS) was completed by utilizing the set of CAAT-box software (7). This program combines the results of different methods: (i)

identification of open reading phases and their tri as a function of their size, (ii) analysis of the probability of being coded by utilizing Genemark software (11), (iii) identification of a start in translation (initiation codon and fixing sequence of the ribosome), (iv) similarity of the proteic sequence deduced with the proteic sequences contained in the sequence banks by utilizing BLASTP software.

The functions of the proteins coded by the coding phases identified were predicted by analysis of the research results of similarities in the non-redundant bank of the NCBI (http://www.ncbi.nlm.nih.gov/BLAST/) by utilizing BLASTP software (1).

4. Comparison of the genomes – identification of the CDS specific to the strain of *L. pneumophila* Paris strain

The set of proteic sequences deduced from the predicted coding phases each genome was compared to the set of proteic sequences possibly coded by the other genome by utilizing BLASTP software. A threshold of 75% of identity on the totality of the length of the protein was retained for identifying the proteins specific to an isolate. This very high value was kept since it best allows the orthologous genes of the paralogous genes (6) to be discriminated. For the proteic sequences for which the sequence preservation is high (> at 70%) the preservation of the nucleotidic sequences of the genes will also be high and could give a signal in hybridization conditions of low stringency. It will be necessary to take into consideration this eventuality in the analysis of the test result.

- 5. Examples of annotations
- 5.1. Genes specific to L. pneumophila Lens strain. There is no significant similarity between the nucleotidic sequence of the gene of L. pneumophila Lens strain and the genome of L. pneumophila Paris strain.

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ID of L. pneumophila Lens strain gene	ID of L. pneumophila Lens strain gene (best score)	% of identity of proteic sequences	% of identity of nucleotidic sequences
2795.1	-	-	-
560.1	-	•	-
116.1	- .		-
3866.1	2661.2	26%	-
2141.1	152.3	24%	not significant

5.2. Genes common to the two strains for which the similarity (identity) of the deduced proteic sequences is less than 75% and value of the similarity at the nucleotide level.

ID of L. pneumophila Lens strain gene	ID of L. pneumophila Paris strain gene (best score)	% of identity of proteic sequences	% of identity of nucleotidic sequences
2518.1	5987.2	42%	32%
3820.1	3661.4	42%	15%

5.4. Genes common to L. pneumophila Lens strain and Paris strain.

ID of L. pneumophila Lens gene	ID of L. pneumophila Paris gene (best score)	% of identity of proteic sequences	% of identity of nucleotidic sequences
795.1	3838.3	99%	98%
2457.1	3282.3	100%	98%

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- 3. Birnboim, H. C. 1983. A rapid alkaline extraction method for the isolation of plasmid DNA. *Methods Enzymol*. 100:243-255.
 - 4. Buchrieser, C., C. Rusniok, L. Frangeul, E. Couve, A. Billault, F. Kunst, E. Carniel, and P. Glaser. 1999. The 102 kb locus of *Yersinia pestis*: sequence analysis and comparison of selected regions among different *Yersinia pestis* and *Yersinia pseudotuberculosis* strains. *Infect. Immun.* 67:4851-4861.
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- 7. Frangeul L, Glaser P, Rusniok C, Buchrieser C, Duchaud E, Dehoux P and Kunst F. 2004. CAAT-Box, Contigs-Assembly and Annotation tool-box for genome sequencing projects. *Bioinformatics*. in press.
- 8. Gordon, D., C. Abajian, and P. Green. 1998. Consed: a graphical tool for sequence finishing. *Genome Res.* 8:195-202.
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Example 8: Proof in the genome of Legionella pneumophila of the exploitation of functions of the host cell and of the high genomic plasticity

Legionella pneumophila, the causal agent of legionnaires disease, replicates in the form of an intracellular parasite of the amoeba and persists in the environment in the form of a free living microbe. Analyzed here are the complete genomic sequences of L. pneumophila Paris (3 503 610 bp; 3 077 genes), an endemic strain predominant in France, and L. pneumophila Lens (3 345 687 bp; 2932 genes), an epidemic strain responsible for a major epidemic in France. A striking characteristic of the genome of L. pneumophila is its plasticity. Three different plasmids were identified, and ~13 % of each genome is different to the other strain. The Paris strain codes for a unique secretion system of type V, and its secretion system Lvh of type IV is coded by a region of 36 kb which can be either carried on a multicopy plasmid or be integrated into the chromosome. The genetic mobility can be a mechanism which increases the multiplicity of L. pneumophila. A large number of genes codes for proteins or patterns of eukaryotic type provided to modulate the functions of the host cell to the advantage of the pathogen, comprising the repeated sequences of tetratrico peptide, ankyrin, F box, serinthreonin kinase proteins, apyrases and a sphingosine-1-phosphate lyase. Therefore, the genome reflects the history and the lifestyle of L. pneumophila, a human pathogen of the macrophages which has co-evolved with soft-water amoeba.

L. pneumophila is the causal agent of the legionellosis, an atypical pneumonia, which can be fatal if it is not rapidly treated. This Gram-negative facultative

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intracellular pathogen can at the same time adapt to the aquatic environment and to the intracellular medium of the phagocytary cells of the human host². When inhaled in contaminated aerosols, *L. pneumophila* can reach the alveolae of the lungs where they swamped by macrophages. By opposition to the majority of bacteria, which are destroyed, *L. pneumophila* can multiply within the phagosome of the macrophage and in the end kill off the macrophage, resulting in legionellosis³.

L. pneumophila and other legionelles are inhabitants of natural aquatic biotopes and artificial aqueous systems, such as the cooling towers of air conditioners³. Legionelles were detected by culture in soft-water environments at 40 % and by PCR in soft-water environments up to 80 %, where the bacteria are known to survive and replicate by intracellular means in free living protozoons, often within aquatic biological films⁴. Its capacity for exploiting the base cellular mechanisms of a large spectrum of protozoic eukaryotic hosts likewise allows legionelles to infect human cells⁵. In fact, it was shown that the capacity of L. pneumophila to multiply by intracellular means in amoeba contributes to the disease, even though little is known of things about the mechanisms governing the host-microbe interactions. Inversely, the biphasic life cycle of L. pneumophila, the changing of replicating parasitic cells into transmissible extracellular forms and the complex regulatory network, which governs these changes, are understood in part⁶.

The Legionella genre comprises 48 species, but more than 90 % of the cases of clinical legionellosis are caused by L. pneumophila even more arresting, up to 84 % are caused by the serogroup 1 of L. pneumophila⁷. We have determined the complete genomic sequences of two clinical isolates of the serogroup 1, the Paris and Lens strains, to provide knowledge of the genetic characteristics of L. pneumophila, and to identify the properties which were selected in niches specific to the pathogenicity and the life cycle of L. pneumophila. The Paris strain is the only endemic strain known to date, accounting for 12.7 % of the cases of legionellosis in France and for 33 % of those occurring in the Paris region⁸. It is associated with the nosocomial and community diseases occurring in the form of epidemics or sporadic cases. From November 2003 to January 2004 the Lens strain caused an epidemic of 86 cases resulting in 17 deaths in the north of France, suggesting that it is particularly efficacious for causing the disease in humans. The genomic comparatives of an endemic isolate and of an epidemic isolate supplying the bases for comprehending the specificity of strain, and can give indices for the particular adaptability and stability of the Paris strain.

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Results

General characteristics

The Paris strain and the Lens strain of *L. pneumophila* each contain a circular chromosome of ~3 503 610 bp and ~3 345 687 bp, respectively, with an average G+C content of 38 % (Table XXII, Figure 1, Gen-Bank/EMBL access numbers CR628336, CR628337). *L. pneumophila* Paris separates a plasmid of 131 885 bp and the Lens strain contains another plasmid of 59 832 bp (Gen-Bank/EMBL access numbers CR628338, CR628339. In the chromosome of *L. pneumophila* Paris, 3077 genes were identified and 2 932 in that of *L. pneumophila* Lens. No function was able to be predicted for 42.1 % (1354) of the genes of *L. pneumophila* Paris and 44.1 % (1320) of the genes of *L. pneumophila* Lens, a proportion similar to that found in the majority of the other bacterial genomic sequences. A high proportion of the genes provided (21 % for the Paris strain, 20.4 % for the Lens strain) is unique to the *Legionella* genre and can thus code the specific functions of *Legionella*.

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Exploitation and modulation of the functions of the host cell

A fascinating question is to know how Legionella to decompose the functions of the host to enter, survive, replicate and evade amoeba or alveolar macrophages. Within its genome L. pneumophila codes for an abundance of proteins of eukaryotic type. In effect, 30 have the highest similarity with eukaryotic proteins (Table XXIII) and 32 genes code for proteins with eukaryotic domains implied in protein–protein interactions (Table XXIV). We reveal here proteins provided for diverting eukaryotic regulatory paths or for being secreted in eukaryotic cells, making strong candidates of them for directing the invasion of Legionella, for traveling in the host cell, or modulating or being subtracted from functions of the host cell.

The repeated sequences of tetratrico peptide (TPR) are repeated patterns of the 34 degenerated amino-acids present in networks in tandem of 3 to 16 patterns, which form hooks for facilitating the protein-protein interactions. The TPR proteins contribute to control of the cellular cycle, to repression of transcription, to response to stress, to inhibition of the protein kinase, to the transport of mitochondrial and peroxisomal protein and to neurogenesis⁹. The Sel-1 repeated sequences represent a sub-family of TPR sequences. In *L. pneumophila* five proteins containing Sel-1 domains were identified. Two of them (EnhC and LidL) were previously implied in interaction with the host cells or in precocious signaling of events which regulate the scheduling

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decisions of L. pneumophila in the macrophages^{10,11}. As a consequence, the three newly identified proteins are likewise in al likelihood implied in the host–pathogen interactions (Table XXIV).

After internalizing, L. pneumophila manipulates the endosome-lysosome degradation path of the host for surviving and replicating within a vacuole derived from the endoplasmic reticulum (ER). A protein of L. pneumophila, RalF, thought to contribute to recruitment of the ER contains a eukaryotic domain Sec 7. RalF, a substrate of the secretion system of type IV is required by the regulatory protein ARF for associating with the phagosomes of L. pneumophila¹². The two strains of L. pneumophila code for three serin/threonin kinase proteins of eukaryotic type (STPK) (Table XXIV). The multisequence comparisons of the domains of the kinase from the Paris strain of L. pneumophila and other prokaryotic and eukaryotic STPK, have revealed that Lpp2626 and Lpp1439 of L. pneumophila aggregate in the group of eukaryotic STPK, close to the STPK originating from Entamoeba histolytica (Figure 2). Mycobacterium tuberculosis, which as L. pneumophila blocks the phagosome-lysosome fusion, produces eleven STPK of eukaryotic type¹³. In particular, the STPK PknG of M. tuberculosis is an inhibitor of the phagosome-lysosome fusion and a promoter of intracellular survival¹⁴. The STPK domain of Lpp0267/Lpl0262 is related to the PknG and to the STPK YpkA of Y. pseudotuberculosis (Figure 2), an enzyme which is translocated in the eukaryotic cells where it corrects the defenses of the host by interfering with the transduction paths of the eukaryotic signal¹⁵. This suggests that the STPK of L. pneumophila can likewise modulate the transduction mechanisms of the eukaryotic signal and can modify the routing paths of the host cell.

Twenty proteins contain ankyrin domains (Table XXIV), sequences repeated in tandem of around 33 amino acids which represent one of the modular interaction protein-protein patterns, the most current being eukaryotic. To date, the only prokaryotic genomes known for coding large families of proteins of the ankyrin domain are *Coxiella burnetii* and *Wolbachia pipientis*, which code 13 and 23 elements, respectively^{16,17}. Similar to *L. pneumophila*, *C. burnetii* is an intracellular pathogen which is extremely well-adapted to life inside the eukaryotic phagolysosome, and *W. pipientis* is an "endosymbiont" parasite living in the reproductive cells of a large variety of arthropods. Therefore, the ankyrin domains can be implied in a common microbial mechanism for manipulating the physiology of the host cell. A possible function of the proteins containing the repeated ankyrin sequences of *L. pneumophila* is to modify the

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interactions with the host cytoskeleton, given that it is thought that numerous eukaryotic ankyrin proteins are binders between the membranous proteins and the cytoskeleton¹⁸ and are important for targeting proteins towards the plasmic membrane or towards the endoplasmic reticulum. The ankyrin domains are likewise compounds of transcription regulators, suggesting that they can influence the expression of the genes of the host cell as proposed for *Ehrlichia phagocytophila*¹⁹. In effect, one of the repeated ankyrin proteins of *L. pneumophila* (Lpp3991/Lpl0559) likewise contains a eukaryotic SET domain, known for binding the chromatin host and for influencing the expression of the genes of the host cell²⁰. In opposition to the ankyrin proteins of *W. pipientis*, none of the repeated ankyrin proteins of *L. pneumophila* contains a peptide signal¹⁷. Instead of this, certain of them could be secreted by means of the secretion system of type IV, a way which is independent of the typical targeting signals.

The final stages in the intracellular life cycle of L. pneumophila are to kill and escape from its host cell, a mechanism which is still not understood. One class of proteins which can affect the control of the division of the host cell (Lpp2082/Lpl2072, Lpp2486 and Lpp0233/Lpl10234) separates eukaryotic F box domains, sites of proteinprotein interactions. Generally, Lpl2072, Lpp2486 and Lpp0233/Lpl10234) separates eukaryotic are associated with other interaction domains²¹. Similarly, two of the identified F box proteins are associated with a repeated ankyrin sequence or a doublespooling pattern, respectively (Table XXIV). The proteins of the F box, assembled in SCF ubiquitin-ligase complexes, determine which substrates are going to be targets for ubiquitination and subsequent proteolysis by the proteasome. Given that the targeted substrates comprise promoters and inhibitors of the cellular cycle as well as transduction compounds of the signal²², the F box proteins can regulate the division and cellular differentiation. To our knowledge, the only protein of prokaryotic F box described is VirF of Agrobacterium tumefaciens, a protein which, it is thought, interacts with the proteins of the host by means of its F box domain to target it for proteolysis²³. Another pattern implied in the eukaryotic ubiquitination is the U box pattern. The protein Lpp2887, present in the Paris strain but not in the Lens strain, contains such a pattern. It is apparently the first recognized in a prokaryotic organism.

Additional proteins of eukaryotic type identified in the genome of the two strains of *L. pneumophila* are sphingosine-1-phosphate lyase (Lpp2128/Lpl2102) and two secreted apyrases (Lpp1033, Lpp1880/Lpl1000, Lpl1869), suggesting that *L. pneumophila* modulates the cycle of the host cell to its advantage. The broadly

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expressed lyase sphingosine-1-phosphate enzyme catalyses the essentially irreversible splitting of the sphingosine-1-phosphate signaling molecule, a product of the degradation of sphingomyeline which regulates cellular proliferation and cellular death in the eukaryote. In effect, the overexpression of the sphingosine-1-phosphate lyase can induce apoptosis in the eukaryote, identifying this enzyme as a double modulator of the metabolism of the sphingosine-1-phosphate and of the ceramid, as well as a regulator of the decisions of cellular direction²⁴. In addition, the sphingosine-phosphate plays a central rôle in the development of the Dictyostelium discoideum amoeba, given that interruption to the gene results in aberrant distribution of the actin, an abnormal morphogenetic phenotype and a viability occurring during the stationary phase²⁵. The two apyrases of L. pneumophila are the only ones identified in the prokaryote. The family of apyrase protein comprises enzymes capable of splitting the tri- and diphosphates nucleotides (NTP and NDP) in a calcium- or magnesium-dependent manner. The apyrase was isolated in the autophagy²⁶ vacuole suggesting that these two proteins influence the destiny of the phagosome of L. pneumophila in diminishing the concentration of NTP and NDP during parasitism of the cell.

246 proteins (7.6 %) in the Paris strain and 231 proteins (7.7 %) in the Lens strain have likewise been identified with double-spooling domains provided (CC), many of which likewise show slight similarities with eukaryotic proteins. The CC domains facilitate the protein-protein interactions either for the multimerisation of protein or the macromolecular recognition. Therefore, double-spooling domains can target proteins at the appropriate locale in the eukaryotic host. Interestingly, all the new substrates (SidA-H, SdeC) of the secretion system of type IV identified by Luo and Isberg²⁷ as well as LidA, LepA and LepB contain double-spooling domains.

To affect the eukaryotic cell, *L. pneumophila* must translocate these proteins of eukaryotic type to the host cytoplasm. As a consequence these proteins are candidate substrates for the secretion system of type IV, as shown for VirF in *A. tumefaciens*²⁸ or RalF in *L. pneumophila*¹².

Secretion system

At the centre of the pathogenesis of *L. pneumophila* are the loci dot/icm, which together direct the assembling of secretion apparatus of type IV. Even though the two strains contain the complete loci dot/icm, their sequences have variations. In effect, the comparison of the sequence of dot/icm genes of 18 different strains of *L. pneumophila*

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has identified a wide range of variations in sequence, and has placed the strains in seven phylogenetic different groups²⁹. However, no correlation with the virulence is apparent.

A novel factor of putative virulence of L. pneumophila, limited to the Paris strain, is a provided auto-transporter protein. Lpp0779 contains several marks of secretion systems of type V comprising a peptide of N-terminal head for secretion across the internal membrane and a specialized C-terminal domain which forms a pore in the other membrane through which the domain clone passes to the surface of the cell³⁰. Its domain clone is a compound of known repeated sequences of haemagglutinin to be implied in the cell-cell aggregation and extremely similar to those of the autotransporter of AIDA-I and Ag43 of Escherichia coli, two proteins implied in the virulence. The bacterial surface protein AIDA-I facilitates adherence to the mammal cells³¹, while Ag43 imparts not only a low level of adhesion to certain mammal cells, but likewise facilitates auto-aggregation which is important for the formation of biological film³². In a similar way, the auto-transporter of L. pneumophila can be implied in adhesion to the host cell and the formation of biological film. In opposition to AIDA-I and Ag43, the auto-transporter of L. pneumophila does not have an RGD pattern implied in the bond with the human integrins, and can thus have another interaction domain. The auto-transporter was acquired in the same way by horizontal gene transfer as suggested from its numerous IS upstream and from GC contents of 41 % which exceed the average of the genome of 38 %. Studies of the distribution of this gene in clinical and environmental strains of L. pneumophila in combination with the study of its function must provide knowledge of its importance.

In addition, the two strains of *L. pneumophila* contain a secretion method by translocation (Tat) with combined arginine (TatAB and TatC) and completes the secretion systems of type I and II. The system of type II coded by the genes *lspA*, *lspD-J* is required for the secretion of several enzymes such as lipase A and B (Lpp0533, Lpp1159/Lpl0509, Lpl1164), phosphatase acid Map and SurE (Lpp1120, Lpp1245/Lpl1124, lpl1245), lysophospholipase A (Lpp2291/Lpl2264) and phospholipase PlaB (Lpp1568/Lpl422), proteins which are all present in the two strains.

<u>Metabolism</u>

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The metabolic paths utilized by *L. pneumophila* for multiplying inside the eukaryotic cells are not known. The bacteria seems to prefer the proteic substrates, given that a large number of absorption and degradation systems of oligopeptide and

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amino acid are coded in the genome. In particular, apart from the elastase homologue ProA of *Pseudomonas aeruginosa* (Lpp0532/lpl0508), three secreted paralogue metalloproteases as well as 46 additional peptidases are present.

By way of opposition, systems for the absorption of sugar are rare, even though the complete ways of Embden-Meyerhof and Entner-Doudoroff are present. In the two strains of L pneumophila no absorption system of type PTS was identified. However, certain of the transport systems of type 55 ABC can be implied in the absorption of sugar, given that the bacteria has some systems for the degradation of complex sugars, such as trehalase, polysaccharide deacetylase, glucoamylase of type eukaryotic (Lpp0489/Lpl0465), β -hexosaminidase and chitinases (Lpp1117/Lpl1121). The two strains code for proteins highly homologous to transporters of glycerol phosphate ABC (Lpp1696, Lpp1695, Lpp1694/Lpl1695, Lpl1694, Lpl1693), and for a hexose phosphate transporter (Lpp2623/Lpl2474), which can be important during intracellular growth. We have likewise identified several enzymes probably implied in the utilization of mesoinositol, which can interfere with the signaling of the host cell facilitated by this intracellular messenger.

L. pneumophila is provided for coding for an extensive aerobic respiratory chain constituted **NADH** deshydrogenase, cytochrome-dependent succinate deshydrogenase, ubiquinol-cytochrome reductase and four terminal oxydases, which guarantee the capacity to adapt to changing oxygen tensions (one cytochrome aa3, two quinol-cytochromes of type bd and one quinol cytochrome oxidase of type o). The latter oxidase is absent in the Lens strain. Systems implied in anaerobic respiration are apparently absent in all the strains. The two genomes code for an ATP synthase of type F_0F_1 typical of γ -proteobacteria, whereas the Paris strain codes for a second ATP synthase similar to the non-characterized systems of archeobacteria and bacteria marines. L. pneumophila likewise codes for at least four sodium/proton anti-carriers (Lpp1464, Lpp2448, Lpp0868, Lpp0667/Lpl1519, Lpl2304, Lpl0839, Lpl0651), which modulate presumably the proton and sodium gradient across the cytoplasmic membrane. As a consequence, a sodium motor force can be utilized for the cellular activities. In this respect, the presence of a compound of polar flagellar motors of type sodium, MotY, thus two significantly different aggregates of gene MotA-MotB, leads to the prediction that mobility can be activated by the sodium motor forces as well as the proton forces. One particular characteristic of L. pneumophila is differentiation in a mature intracellular form which accumulates inclusions of poly-hydroxybutyrate in the form of

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carbon and energy reserve. As a consequence, the Paris strain codes for four paralogue poly-hydroxybutyrate synthases and the Lens strain codes for three paralogue poly-hydroxybutyrate synthases (Lpp2323, Lpp2038, Lpp2214, Lpp0650/Lpl1055a-b, Lpl2186, Lpl0634).

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Physiological adaptation and regulation of the gene

In accord with this intracellular life style the regulatory repertoire is rather small. Analysis of the genome has identified 92 transcription regulators (79 in the Lens strain), which represent only 3.0 % of the genes provided. *L. pneumophila* codes for six sigma putative factors, the homologous of *rpoD*, *rpoH*, *rpoS*, *rpoN*, *fliA* and the sigma factor *rpoE* of type ECF. The number of systems with two compounds (13 histidine kinases and 14 response regulators) is likewise low.

The most abundant class of regulators belongs to the GGDEF/EAL (23) family. Present in numerous bacteria, comprising *Vibrio cholerae* (41), *P. aeruginosa* (33), *Wolinella succinogenes* (26), and *E. coli* (19), these regulators contain two subdomains, GGDEF and EAL. Of the 23 regulators identified, 10 separate only one GGDEF domain, 3 in the Paris strain and 2 in the Lens strain contain an EAL domain, and 10 in the Paris strain and 11 in the Lens strain present a combination of the two. The rôle of these regulators in *L. pneumophila* is unknown, but in other bacteria these regulators play a rôle in aggregation, the formation of biological film or mobility by muscular contraction.

In *L. pneumophila*, the cyclic AMP can likewise translate the cellular signals given that the genome codes for five adenylate cyclases of class III (Lpp1446, Lpp1131, Lpp1704, Lpp1277, Lpp0730/Lpl1538, Lpl1135, Lpl1703, Lpl1276, Lpl0710). In *P. aeruginosa*, CyaB, an adenylate cyclase of class III, is implied in the regulation of the virulence of gene 33. However, *L. pneumophila* does not contain the orthologue of Vfr, the dependent AMPc regulator of *P. aeruginosa*, but it codes for five proteins with AMPc bonding patterns (Lpp3069, Lpp1482, Lpp2063, Lpp0611, Lpp2777/Lpl2926, Lpl1501, Lpl2053, Lpl0592a-b-c, Lpl2648). As for *P. aeruginosa*, these adenylate cyclases of class III can comprise environmental signals extending from the nutritional content of the surrounding medium to the presence of host cells and can control the expression of the virulence of the gene in consequence.

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Heightened plasticity of the genomes of L. pneumophila

The two genomes exhibit an astonishingly high plasticity and a diversity of the genome. Comparison of the chromosomes has identified a preserved skeleton of 2664 genes but 280 and 428 genes (10 and 14 %) are specific to the strain (Figure 3). Given that the two strains analyzed belong to the same species and the same serogroup, this diversity was unexpected. For example the comparison of the genomes of two strains of Salmonella enterica of serotype Typhi identifies only 2 % for each of the genes specific to the strain³⁴. The specific genetic equipment of the Paris strain contains a certain number of regulators (three homologous of CsrA, 13 transcription regulators), of additional repeated ankyrin sequences and proteins of type eukaryotic (Table XXIII and Table XXIV) as well as several restriction modification genes (modification methylases of the DNA, endonucleases), which can explain the low competence (personal observation) and the high genomic stability⁸ of the Paris strain. The Lens strain contains fewer specific regulators (4) and four specific proteins with eukaryotic domains (Table XXIII), two of which are repeated sequences of ankyrin proteins, suggesting that the Paris strain is a particularly well-equipped strain.

The genomes of L. pneumophila have undergone rearrangements of multiple genomes. The important synteny in the genome between the Paris and Lens strain is interrupted by inversion of 260 kb, insertion of 130 kb in the Paris strain (or deletion in the Lens strain) and by deletions and smaller multiple insertions. The fragment of 130 kb is flanked by an ARnt gene and codes for a putative integrase, suggesting a structure similar to the islets of pathogenicity of the enterobacteria. It contains an ATP synthase, chemiosmotic flow systems (cebABC, cecABC) and the genes cadA1, ctpA, copA1, copA2 coding for ATP-dependent flow pumps, as was proven to be induced in the macrophages³⁵ and separate the prpA-lvrABC gene aggregate, present within a pathogenicity islet of 65 kb in the Philadelphia strain³⁶. With the exception of abovementioned genes, this pathogenicity islet is absent from the Paris strain and from the Lens strain. However, the corresponding chromosomal site in the Paris strain is the insertion site of an integrative plasmid discussed below. Therefore, these two regions can be hot points for genomic rearrangements. Genomic variation is likewise evident from its network of mobile elements comprising ten integrases, 58 insertion sequences (34 complete and 24 truncated) thus as proteins relevant to phages. In addition, the genomes contain a large number of repeated sequences organized in the form of repeated inverse sequences, which recall the ERIC sequences of the enterobacteria.

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These LeRIC (Repeated Intergenic Consensus of Legionella) fall into 7 classes present in numerous copies (for example 80, 18, 18, 25, 9, 9 and 6 in the Paris strain).

L. pneumophila Paris and Lens contain lvh, a region which codes for a second secretion system of type IV previously characterized in the Philadelphia strain³⁷. One interesting observation is that the lvh region of L. pneumophila Paris is coded on a region of 36 kb which exists either integrated into the chromosome or excised in the form of a multicopy plasmid (unpublished data). This pattern is similar to that described for the unstable element of 30 kb of the Olda strain, which is possibly phage-derived and is implied in the phase variation³⁸. The GC content of the lvh region (43 %) is different to that of the remainder of the chromosome (38 %) and it contains certain genes related to phages, suggesting possible phagic origin. However, the exact excision and integration mode is still not understood. An attractive hypothesis is that the integration and excision of particular regions of the chromosome is a mechanism specific to L. pneumophila for boosting versatility.

The second plasmid of the Paris strain (132 kb) comprises known virulence factors, mobile genetic elements and genes of antibiotic resistance. The regulator system with two lrpR-lskS compounds present on this plasmid was found on a plasmid of *Legionella longbeachae* implied in the virulence of these species³⁹. Heightened preservation (93 to 98 % of protein identity) of the six gene sequences on the plasmid of 135 kb along *L. longbeachae* can indicate a recent horizontal transfer between *L. pneumophila* and *L. longbeachae*. *L. pneumophila* Lens contains a plasmid of 60 kb which codes for several proteins homologous to the transfer region of the F plasmid of *E. coli*. All three plasmids of the Paris and Lens strain code for a paralogue of CsrA, a repressor of the transmission traits and activator of replication⁴⁰.

Although the rôle of the plasmids in the virulence of L. pneumophila has yet to be determined, the correlation between strains containing a plasmid and the virulence in a mouse model was described⁴¹. In addition, L. pneumophila strains with plasmids seem to persist longer in the environment than those strains not having plasmids⁴². The identification in the clinical isolates of the plasmids coding for factors of putative virulence is another indication of the importance of the plasmids for the pathogenicity of Legionella.

The genomes of *L. pneumophila* display a plasticity likewise at the gene level. The loci *enh*, implied in the entry in the host cells⁴³, are present in the Paris and Lens strains. One of the proteins coded by these loci is RtxA, which contributes to entry,

adherence, cytotoxicity, pore formation⁴³ and intracellular routing in the amoeba⁴⁴. Unlike the AA100¹⁰ strain in the Paris strain, rtxA is fused with arpB and a second protein with about 30 sequences repeated in tandem extremely preserved de 549 bp. A similar structure is coded by the Lens strain, however we have identified two patterns within the repeated region, the two being different to that of the Paris strain. However, the number of repetitions seems to be the same (Figure 4). It is possible that the variations in number and sequence of the repeated sequences contribute to the multiplicity and likewise to the virulence of *L. pneumophila*.

In accordance with the relative plasticity in their genome, the strains of L. pneumophila code for organelles of type IV bacterial pili, which are required for natural competence for the transformation of the DNA⁴⁵. The organization of the genes coding for type IV bacterial pili is similar to that of P. aeruginosa where they are crucial for bacterial adherence and colonization of the mucosal surfaces and for mobility by muscular contraction. Another mechanism in L. pneumophila contributing to the plasticity of the genome is conjugated transfer facilitated by the secretion of type IV of plasmids⁴⁶ and chromosomal DNA⁴⁷.

Conclusion

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Analysis of the sequences of the genome of the clinical of L. pneumophila Paris and Lens strains and its comparison identifies L. pneumophila as an extremely versatile organism, which demonstrates a plasticity and an extensive genomic diversity. The excision and integration of plasmids or genes can be a mechanism which L. pneumophila exploits for adapting to different environments. Its large cohort of proteins of eukaryotic type is provided for manipulating the host cell to the advantage of the pathogen (Figure 5). Eucaryotic proteins of putative origin have likewise been identified in other intracellular pathogens, comprising Coxiella, Wolinella, Agrobacterium, Mycobacterium and Ehrlichia, but currently L. pneumophila sequences them as prokaryotic with the greatest variety of proteins of eukaryotic type. Presumably, during its co-evolution with free living amoeba, the L. pneumophila pathogen acquires DNA by horizontal transfer from its host or by convergent evolution. These proteins can then likewise contribute to the infection of human macrophages. By being based on the genomic sequences future comparative and functional studies are going to enable survival tactics of intracellular parasites to be defined, and to identify the special attributes of endemic and epidemic L. pneumophila. To combat the menace of L.

pneumophila resistant to chemical products widely utilized for decontamining public water systems, comprising hospitals, the genomic sequence can stimulate the identification of targets for novel active biocides against L. pneumophila.

5 Methods

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Preparation and sequencing of the DNA. Paris and Lens strains of L. pneumophila were cultivated on agar BCYE at 37°C over 3 days and the chromosomal DNA was isolated by utilizing standard protocols. The cloning, sequencing and assembling were completed as described previously⁴⁸. For the two genomes, two libraries (inserts of 1-2 kb and 2-3 kb) were generated by random mechanical chiseling of the genomic DNA and cloning in pcDNA-2,1 (Invitrogen). A hook was obtained by terminal sequencing of clones from a BAC library constructed as described previously⁴⁹ by utilizing pIndigoBac (Epicentre) as vector. For the Paris strain of L. pneumophila an insert library of average size (8-10 kb) was constructed in the low-number vector of copies pSYX34. The purification Plasmide DNA was produced either with Montage Plasmide Miniprep96 Kit (Millipore) or by utilizing the DNA sequencing matrix amplification kit TempliPhi (Amersham Biosciences). The sequencing reactions were created by utilizing the sequencing reactions kit ABI PRISM BigDye Terminator and an analyzer 3700 or a 3730 Xl Genetique Analyzer (Applied Biosystems). 47,200 sequences for the Paris strain of L. pneumophila, and 47,231 sequences for the Lens strain each originating from four libraries were obtained and assembled and finished as described previously 48.

Annotation and analysis. The definition and annotation of the coding sequences (CDS) is as was described previously⁴⁸ by utilizing the Boîte CAAT 50 software. All the CDS provided were checked visually. The predictions on function were based on preferred BLASTp similarity and on analysis of patterns by utilizing the PFAM, Prosite and SMART databases. We have identified orthologous genes by better BDERNIÈRE reciprocal correlation and FASTA comparisons. For identification of the double-spooling domains the publicly available software PairCoil and Coilscan were utilized. The pseudogenes have one or more mutations which prevent complete translation. Repetitive sequences of DNA were identified by BLASTN comparisons of the intergenic regions and of the complete genome. MFOLD software was utilized to predict the folding of the single sheet of DNA molecules.

URL. The sequence and annotation of the two genomes of *L. pneumophila* are at http://genolist.pasteur.fr/LegioList. For annotation and analysis we used PairCoil http://paircoil.lcs.mit.edu/cgi-bin/paircoil and Coilscan http://www.biology.wustl.edu/gcg/coilscan.html

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Table XXII: General characteristics of the two Legionella pneumophila genomes

	L. pneumophila Paris	L. pneumophila Lens
Size of the chromosome (bp)	3 503 610	3 345 687
G+Ct content	38.3% (37.4%)	38.4% (38.4%)
G+C protein-coding genes content	39.1% (37.9%)	39.4 %(39.1%)
Total number of protein-coding genes	3076 (141)	2931 (57)
Average length (codons) of protein-coding genes	331	333
Number of operons rRNA (16S-23S-5S)	3	3
Number of genes tRNA	43	43
Coding in percentage	87.9% (92%)	88% (83.7%)
Plasmid	1 (131.9 kb)	1 (59.8 kb)
Number of genes specific to the line	428 (125)	280 (44)
Number of orthologous genes	2664	2664

¹kilobase pairs

5 Table XXIII: Proteins having the greatest similarity with eukaryotic proteins

L. pneumophila Paris	Produit provided	L.p.Lens	G-C	Percentage of protein identity
lpp1647	purC	lpl1640	38%	61% over the entire length (AAR06292.1 Nicotania tabacum)
lpp0702	exoA exodeoxyribonuclease III	lpl0684	39%	58% over the entire length (EAA20230.1 Plasmodium yoelii yoelii)
lpp0321	Precursor protein bond to RNA	-	34%	45% of 50% of the protein (AAL07519 Solanum tubeosum)
lpp1157	pyruvate decarboxylase	lpl1162	39%	50% over the entire length (AAB16855.1 Arabidopsis thaliana)
lpp1522	protein de biosynthesis de thiamine NMT-1	lpl1461	38%	49% over the entire length (AAC64375.1 Botryotinia fuckeliana)
lpp2832	nuoE NADH dehydrogenase I chain E	lpl2701	38%	49% of 82% of the protein (BAA25988.1 Homo sapiens)
plpp0050	protein with hypersensitive induced response	-	36%	48% over the entire length (AAN17462.1 Hordeum vulgare subsp. Vulgare)
lpp0634	hypothetical protein	lpl0618	39%	48% over the entire length (XP_306643.1 Anopheles gambiae)
lpp0965	protease	lp10935	39%	45% over the entire length (NP_189431.2 Arabidopsis thaliana)
lpp2748	phytanoyl-coA dioxygenase	lpl2621	36%	44% over the entire length (XP_372144.1 Homo sapiens)
lpp2128	sphingosine-1-phosphate lyase	lp12102	41%	36% over the entire length (NP_7.75139.1 Rattus norvegicus)

lpp0489	glucoamylase	lpl0465	39%	32% over the entire length (P42042 Arxula adeninivorans)
lpp0955	cytokinin oxydase	lpl0925	39%	32% over the entire length (NP_484368.1 Nostoc sp.)
lpp0578	phytanoyl coA dioxygenase	lpl0554	36%	31% over the entire length (EAA70100.1 Gibberella zeae)
lpp0379	hypothetical protein	Lpl0354	39%	31% over the entire length (CAD21525.1 Taenia solium)
lpp1033	ectonucleoside triphosphate diphosphohydrolase (apyrase)	lpl1000	40%	25% over the entire length - nucleoside phosphatase signature (Q9MYU4 Sus scrofa)
lpp2923	6-pyruvoyl-tetrahydropterin synthase	lp12777	34%	26% over the entire length (NP_703938.1 Plasmodium falciparum)
lpp3071	zinc metalloproteinase	lpl2927	38%	38% over the entire length (AAF56122.1 Drosophila melanogaster)
lpp2134	methyltransferase bonded to SAM	lpl2109	35%	24% over the entire length (BAC98835.1 Bombyx mori)
lpp1880	ectonucleoside triphosphate diphosphohydrolase (apyrase)	lpl1869	39%	26% over the entire length (CAE70887.1 Caenorhabditis briggsae)
lpp2747	methyltransferase bonded to SAM	lpl2620	35%	33% of 56% of the protein (EAA20288.1 Plasmodium yoelii yoelii)
lpp2468	Cytochrome P450	lpl2326	39%	20% of 75% of the protein (NP_487786.1 <i>Nostoc sp.</i>)
lpp1824	protein bound to the nuclear membrane	-	34%	19% of 40% of the protein (NP_082559.1 Mus musculus)
lpp1665	uracyl DNA glycosylase	lpl1659	36%	27% of 80% of the protein (EAA36774.1 Giardia lamblia)
lpp1959	condensation chromosome type 1	lpl1953	41%	Model of preserved chromosome condensation regulator
lpp0358	hypothetical protein	lpl334	38%	37% on 53% of the protein (EAA20288.1 Plasmodium yoelii yoelii)
lpp1127	Ca2+-ATPase de transport	lp11131	37%	22% on 34% of the protein (AAB81284.1 Paramecium tetraurelia)
lpp1167	uridine kinase	lpl1173	33%	35% on 65% of the protein (AAM09314.2 Dictyostelium discoideum)
lpp2626	domain ser/thr kinase protein	lp12481	32%	domain preserved
lpp1439	serine threonine kinase protein	lp11545	36%	domain preserved

Lpp indicates the coding sequences (CDS) provided from L. pneumophilastrain Paris; lpl indicates the coding sequences (CDS) provided from L. pneumophilastrain Lens, the lines in gray indicate the proteins which are likewise mentioned in Table 2B in terms of their preserved eukaryotic domains; the access numbers to the proteins are indicated between parentheses

<u>Table XXIV</u>: coding domains of L. pneumophila preferably protein found in eukaryotic proteins

L. pneumophila	L. pneumophila	identified unit	Content G-C	putative function
Paris	Lens			
enhC (lpp2692)	enhC (lpl2564)	21 domains sel-1	39%	
lidL(lpp1174)- EnhC paralog lpp1310- EnhC	lidL (lpl1180) - EnhC paralog lpl1307 - EnhC	6 domains sel-1	38%	Invasion and traffic in host cells
paralog lpp2174- EnhC	paralog lpl1303 - EnhC	4 domains sel-1	41%	
paralog	paralog lpl1059 - EnhC	3 domains sel-1	40%	
	paralog	7 domains sel-1	45%	
ralF (lpp1932)	ralF lpl1919	domain sec7	34%_	
lpp0267	lpl0262	domain ser/thr kinase protein	38%	
lpp2626	lpl2481	domain ser/thr kinase protein	32%	Modulation functions of host cells
lpp1439	lpl1545	domain ser/thr kinase protein	36%	
lpp2065	lp2055	ankyrin repetition	37%	
lpp0037	lp10038	ankyrin repetition	38%	
plpp0098	-	ankyrin repetition	37%	
lpp2058	lpl2048	ankyrin repetition	38%	
lpp0750	lpl0732	ankyrin repetition	35%	
lpp2061	lpl2051	ankyrin repetition	39%	
lpp2270	lp12242	ankyrin repetition	34%	
lpp0503	lp10479	ankyrin repetition	38%	
lpp1905	_	ankyrin repetition	35%	
lpp1683	lpl1682	ankyrin repetition + domain SET	33%	
lpp2248	lpl2219	ankyrin repetition	39%	
lpp0202	-	ankyrin repetition	38%	
lpp0469	lpl0445	ankyrin repetition	38%	
lpp2517	lpl2370	ankyrin repetition	36%	
lpp1100	-	ankyrin repetition	48%	
lpp0126	lpl0111	ankyrin repetition	39%	
lpp0356	-	ankyrin repetition	38%	
lpp2522	lpl2375	ankyrin repetition	39%	
lpp0547	lp10523	ankyrin repetition	40%	
-	lpl1681	ankyrin repetition	34%	
-	lpl2344	ankyrin repetition	35%	
-	lpl2058	ankyrin repetition	40%	
lpp2082	lpl2072	domain F-box + ankyrin repetition	36%	
lpp2486	-	domain F-box + superhelice	34%	Control division, evasion host cells
lpp0233	lpl0234	domain F-box	2007	
lpp2887	-	2 domains U-box	35%	
lpp2128	lpp2102	2 domining of box	22/0	
Sphingosine-1- phosphate lyase	Sphingosine-1- phosphate lyase		41%	

Lpp indicates the coding sequences (CDS) provided from L. pneumophilastrain Paris; lpl indicates the coding sequences (CDS) provided from L. pneumophilastrain Lens; the numbers indicate the number of domains identified in a protein; ser/thr indicates threonine serin

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Example 9: Repeated sequences of DNA and secreted enzymes characteristic of Legionella pneumophyla

A) Repeated sequence

A repeated sequence was identified in the genome of the Legionella pneumophila Paris strain then sequenced completely. This sequence (SEO ID 7074) is of 122 bp and is repeated 86 times in the genome of the L. pneumophila Paris strain. The preservation is from 81 to 100 % (0 to 19 mismatch) over the entire length for 53 copies and from 70 to 80 % over a length of at least 100 nucleotides for 33 copies. In the L. pneumophila Lens strain, there are 62 copies whereof the preservation of 29 copies is from 81 % to 95 % over the entire length. We have determined oligonucleotides specific to this sequence for its amplification by PCR. Tests on 15 strains each belonging to one of the serotypes of L. pneumophila (serotypes 1 to 15) have shown that this sequence is present in all serotypes of L. pneumophila. The test of 11 strains belonging to other species of the Legionella genre (L. miedadei, L. dumoffii, L. gormanii, L. longbeachae serogroup 1, L. jordanis, L. anisa, L. erythre, L. rubrilucens, L. quinlivani, L. moravica, L. taurinensis) with these oligonucleotides, thus databank research have shown that this sequence is specific to the Legionella pneumophila species and that it will be able to thus serve as an identifier of the species (see Tables XXV and XXVI and Figures 6 and 7). The high number of copies of this sequence in the genome will enable amelioration of the sensitivity of a PCR test or by hybridization, compared to a present sample in a single copy.

SEQ ID 7074:

AGGACTTACGAAAAACCCCAAGATCAAGGCAAAAAATGTTTTTAATGAGG GAGTTTAGATAAACTAAATAACCGAATTAAAAAACTTTTTTTAACAAAGAG

30 ATTGGGATTTTTCGTAAGTCCT.

This sequence is an interesting target for diagnostics of L. pneumophila diagnostic by PCR or by other methods equivalent to PCR.

Among the primers used for amplification and detection of this repeated sequence, the following couple of primers can especially be cited:

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SEQ ID N° 7075: GAAAAACCCCAAGATCAAGGC and

SEQ ID N° 7076: AGGACTTACGAAAAACCCCAA.

<u>Table XXV</u>: List of the DNA of the 15 reference serogroups of *Legionella pneumophila*

Name	Number labo	n°ATCC
L pneumophila sg1	R1	ATCC 33152
L pneumophila sg3	R2	ATCC 33155
L pneumophila sg2 Togus 1	R6	ATCC 33154
L pneumophila sg4 LosAngeles1	R7	ATCC 33156
L pneumophila sg5 subsp fraseri	R8	ATCC 33216
L pneumophila sg6 Chicago 2	R10	ATCC 33215
L pneumophila sg7 Chicago 8	R12	ATCC 33283
L pneumophila sg8 Concord 3	R13	ATCC 33096
L pneumophila sg9 IN-23-G1-E2	R14	ATCC 35289
L pneumophila sg10 Leiden 1	R15	ATCC 43283
L pneumophila sg11 797-PA-H	R18	ATCC 43130
L pneumophila sg12 570-CO-H	R19	ATCC 43290
L pneumophila sg13 82A3105	R20	ATCC 43736
L pneumophila sg14 1169-MN-H	R21	ATCC 43073
L pneumophila sg15 Lansing3	R22	ATCC 35251

<u>Table XXVI</u>: list of the DNA of the 11 reference species Legionella non pneumophila

ATCC Number	Description	Strain Reference
33218	Tatlockia micdadei Garrity et al. deposited as Legionella micdadei	TATLOCK [CIP 103882; NCTC 11371]
33279	Fluoribacter dumoffii (Brenner et al.) Brown et al. deposited as Legionella dumoffii	NY 23
33297	Fluoribacter gormanii (Morris et al.) Brown et al. deposited as Legionella gormanii	LS-13 [ALLO3]
33462	Legionella longbeachae McKinney et al. serogroup 1	Long Beach 4 [NCTC 11477]
33623	Legionella jordanis Cherry et al.	BL-540
<u>35291</u>	Legionella anisa Gorman et al.	CH-47-C3
<u>35303</u>	Legionella erythra Brenner et al.	SE-32A-C8 [NCTC 11977]
35304	Legionella rubrilucens Brenner et al.	WA-270A-C2 [NCTC 11987]
43830	Legionella quinlivanii Benson et al. serogroup 1	1442-AUS-E [CIP 105272]
43877	Legionella moravica Wilkinson et al	316-86 [CDC 1634-CZK-E; CIP 103883]
700508	Legionella taurinensis Lo Presti et al.	Turin I no 1

B) Enzymes secreted

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- The enzymes secreted and common to the three strains of L. pneumophila (Paris, Lens and Philadelphia) whereof the sequences are identified hereinbelow can be utilized especially as a target in colorimetric tests (or for their being made available) for detection of the presence or not of Legionella in a biological sample.
- the sequence lpp0489 (SEQ ID 4292) which codes for a precursor of glucoamylase (Glucan 1,4-alphā-glucosidase) of eukaryotic cell without homologue in the bacteria;
- the sequence lpp1117 (SEQ ID 6477) which codes for a potential secreted chitinase; and

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- the sequences lpp1033 (SEQ ID 4267) and lpp1880 (SEQ ID 3675) which code for a protein similar to an ectonucleoside triphosphate diphosphohydrolase (apyrase) secreted from a eukaryotic cell.

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<u>Table I</u>: Correspondence of the numbers attributed to the genes of the Paris strain on its contigs with the numbers of SEQ ID identified in the list of sequences and position of nucleic sequences coding these genes on the sequence of these contigs with their putative function, as well as their specificity relative to the Philadelphia strain

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Table II « Best-BlastP »: Putative function of the ORFs identified for the Paris strain

<u>Table III</u>: Correspondence of the numbers attributed to the contigs with the numbers of the SEQ ID identified in the list of sequences

Contig1 SEQ ID No. 1 Contig29 SEQ ID No. 29 Contig2 SEQ ID No. 2 Contig30 SEQ ID No. 30 Contig3 SEQ ID No. 3 Contig31 SEQ ID No. 31 Contig4 SEQ ID No. 4 Contig32 SEQ ID No. 32 Contig5 SEQ ID No. 5 Contig33 SEQ ID No. 33 Contig6 SEQ ID No. 6 Contig34 SEQ ID No. 34 Contig7 SEQ ID No. 7 Contig35 SEQ ID No. 35 Contig8 SEQ ID No. 8 Contig36 SEQ ID No. 36 Contig9 SEQ ID No. 9 Contig37 SEQ ID No. 36 Contig10 SEQ ID No. 10 Contig38 SEQ ID No. 37 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 38 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 38 Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 39 Contig14 SEQ ID No. 13 Contig40 SEQ ID No. 40 Contig14 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 14				
Contig3 SEQ ID No. 3 Contig31 SEQ ID No. 31 Contig4 SEQ ID No. 4 Contig32 SEQ ID No. 32 Contig5 SEQ ID No. 5 Contig33 SEQ ID No. 33 Contig6 SEQ ID No. 6 Contig34 SEQ ID No. 34 Contig7 SEQ ID No. 7 Contig35 SEQ ID No. 35 Contig8 SEQ ID No. 8 Contig36 SEQ ID No. 36 Contig9 SEQ ID No. 9 Contig37 SEQ ID No. 37 Contig10 SEQ ID No. 10 Contig38 SEQ ID No. 37 Contig10 SEQ ID No. 11 Contig39 SEQ ID No. 38 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 39 Contig11 SEQ ID No. 12 Contig40 SEQ ID No. 39 Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig13 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 14 Contig42 SEQ ID No. 42 Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 17 <td></td> <td><u> </u></td> <td></td> <td>·</td>		<u> </u>		·
Contig4 SEQ ID No. 4 Contig32 SEQ ID No. 32 Contig5 SEQ ID No. 5 Contig33 SEQ ID No. 33 Contig6 SEQ ID No. 6 Contig34 SEQ ID No. 34 Contig7 SEQ ID No. 7 Contig35 SEQ ID No. 35 Contig8 SEQ ID No. 8 Contig36 SEQ ID No. 36 Contig9 SEQ ID No. 9 Contig37 SEQ ID No. 37 Contig10 SEQ ID No. 10 Contig38 SEQ ID No. 37 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 38 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 39 Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 39 Contig13 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig14 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 14 Contig42 SEQ ID No. 42 Contig14 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 45 Contig17 SEQ ID No. 18<			Contig30	SEQ ID No. 30
Contig5 SEQ ID No. 5 Contig33 SEQ ID No. 33 Contig6 SEQ ID No. 6 Contig34 SEQ ID No. 34 Contig7 SEQ ID No. 7 Contig35 SEQ ID No. 35 Contig8 SEQ ID No. 8 Contig36 SEQ ID No. 36 Contig9 SEQ ID No. 9 Contig37 SEQ ID No. 37 Contig10 SEQ ID No. 10 Contig38 SEQ ID No. 38 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 39 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 39 Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig13 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 13 Contig41 SEQ ID No. 42 Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 44 Contig17 SEQ ID No. 18 Contig44 SEQ ID No. 45 Contig18 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 2	Contig3	SEQ ID No. 3	Contig31	SEQ ID No. 31
Contig6 SEQ ID No. 6 Contig34 SEQ ID No. 34 Contig7 SEQ ID No. 7 Contig35 SEQ ID No. 35 Contig8 SEQ ID No. 8 Contig36 SEQ ID No. 36 Contig9 SEQ ID No. 9 Contig37 SEQ ID No. 37 Contig10 SEQ ID No. 10 Contig38 SEQ ID No. 38 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 39 Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig13 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 14 Contig42 SEQ ID No. 42 Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 43 Contig17 SEQ ID No. 16 Contig44 SEQ ID No. 44 Contig18 SEQ ID No. 18 Contig44 SEQ ID No. 45 Contig19 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 49 Contig21 SEQ ID No.	Contig4	SEQ ID No. 4	Contig32	SEQ ID No. 32
Contig7 SEQ ID No. 7 Contig35 SEQ ID No. 35 Contig8 SEQ ID No. 8 Contig36 SEQ ID No. 36 Contig9 SEQ ID No. 9 Contig37 SEQ ID No. 37 Contig10 SEQ ID No. 10 Contig38 SEQ ID No. 38 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 39 Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig13 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig13 SEQ ID No. 13 Contig41 SEQ ID No. 40 Contig14 SEQ ID No. 14 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 15 Contig42 SEQ ID No. 42 Contig15 SEQ ID No. 16 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 17 Contig44 SEQ ID No. 44 Contig17 SEQ ID No. 18 Contig44 SEQ ID No. 45 Contig18 SEQ ID No. 19 Contig46 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID N	Contig5	SEQ ID No. 5	Contig33	SEQ ID No. 33
Contig8 SEQ ID No. 8 Contig36 SEQ ID No. 36 Contig9 SEQ ID No. 9 Contig37 SEQ ID No. 37 Contig10 SEQ ID No. 10 Contig38 SEQ ID No. 38 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 39 Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig13 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 14 Contig42 SEQ ID No. 42 Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 43 Contig17 SEQ ID No. 17 Contig44 SEQ ID No. 44 Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 50 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 51 Contig24 SEQ ID	Contig6	SEQ ID No. 6	Contig34	SEQ ID No. 34
Contig9 SEQ ID No. 9 Contig37 SEQ ID No. 37 Contig10 SEQ ID No. 10 Contig38 SEQ ID No. 38 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 39 Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig13 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 14 Contig42 SEQ ID No. 42 Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 44 Contig17 SEQ ID No. 17 Contig45 SEQ ID No. 45 Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ	Contig7	SEQ ID No. 7	Contig35	SEQ ID No. 35
Contig10 SEQ ID No. 10 Contig38 SEQ ID No. 38 Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 39 Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig13 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 14 Contig42 SEQ ID No. 42 Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 44 Contig17 SEQ ID No. 17 Contig45 SEQ ID No. 45 Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 53 Contig25 SE	Contig8	SEQ ID No. 8	Contig36	SEQ ID No. 36
Contig11 SEQ ID No. 11 Contig39 SEQ ID No. 39 Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig13 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 14 Contig42 SEQ ID No. 42 Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 44 Contig17 SEQ ID No. 17 Contig44 SEQ ID No. 45 Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 54 Contig27 SE		SEQ ID No. 9	Contig37	SEQ ID No. 37
Contig12 SEQ ID No. 12 Contig40 SEQ ID No. 40 Contig13 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 14 Contig42 SEQ ID No. 42 Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 44 Contig17 SEQ ID No. 17 Contig45 SEQ ID No. 45 Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 55 Contig27 SE	Contig10	SEQ ID No. 10	Contig38	SEQ ID No. 38
Contig13 SEQ ID No. 13 Contig41 SEQ ID No. 41 Contig14 SEQ ID No. 14 Contig42 SEQ ID No. 42 Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 44 Contig17 SEQ ID No. 17 Contig45 SEQ ID No. 45 Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig11	SEQ ID No. 11	Contig39	SEQ ID No. 39
Contig14 SEQ ID No. 14 Contig42 SEQ ID No. 42 Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 44 Contig17 SEQ ID No. 17 Contig45 SEQ ID No. 45 Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig12	SEQ ID No. 12	Contig40	SEQ ID No. 40
Contig15 SEQ ID No. 15 Contig43 SEQ ID No. 43 Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 44 Contig17 SEQ ID No. 17 Contig45 SEQ ID No. 45 Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig13	SEQ ID No. 13	Contig41	SEQ ID No. 41
Contig16 SEQ ID No. 16 Contig44 SEQ ID No. 44 Contig17 SEQ ID No. 17 Contig45 SEQ ID No. 45 Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig14	SEQ ID No. 14	Contig42	SEQ ID No. 42
Contig17 SEQ ID No. 17 Contig45 SEQ ID No. 45 Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig15	SEQ ID No. 15	Contig43	SEQ ID No. 43
Contig18 SEQ ID No. 18 Contig46 SEQ ID No. 46 Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig16	SEQ ID No. 16	Contig44	SEQ ID No. 44
Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig17	SEQ ID No. 17	Contig45	SEQ ID No. 45
Contig19 SEQ ID No. 19 Contig47 SEQ ID No. 47 Contig20 SEQ ID No. 20 Contig48 SEQ ID No. 48 Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig18	SEQ ID No. 18	Contig46	SEQ ID No. 46
Contig21 SEQ ID No. 21 Contig49 SEQ ID No. 49 Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig19	SEQ ID No. 19		SEQ ID No. 47
Contig22 SEQ ID No. 22 Contig50 SEQ ID No. 50 Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig20	SEQ ID No. 20	Contig48	SEQ ID No. 48
Contig23 SEQ ID No. 23 Contig51 SEQ ID No. 51 Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig21	SEQ ID No. 21	Contig49	SEQ ID No. 49
Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig22	SEQ ID No. 22	Contig50	SEQ ID No. 50
Contig24 SEQ ID No. 24 Contig52 SEQ ID No. 52 Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig23	SEQ ID No. 23	Contig51	SEQ ID No. 51
Contig25 SEQ ID No. 25 Contig53 SEQ ID No. 53 Contig26 SEQ ID No. 26 Contig54 SEQ ID No. 54 Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55		SEQ ID No. 24		
Contig26SEQ ID No. 26Contig54SEQ ID No. 54Contig27SEQ ID No. 27Contig55SEQ ID No. 55	Contig25	SEQ ID No. 25		
Contig27 SEQ ID No. 27 Contig55 SEQ ID No. 55	Contig26	SEQ ID No. 26		
	Contig27	SEQ ID No. 27		
	Contig28	SEQ ID No. 28		

<u>Table IV</u>: Correspondence between the contigs of Legionella pneumophila Philadelphia strain and numbering of their sequence in the list of sequences

Contig of the Philadelphia strain	Seq Id
Contig1	SEQ ID N°3456
Contig2	SEQ ID N°3457
Contig3	SEQ ID N°3458
Contig4	SEQ ID N°3459
Contig5	SEQ ID N°3460
Contig6	SEQ ID N°3461
Contig7	SEQ ID N°3462
Contig8	SEQ ID N°3463
Contig9	SEQ ID N°3464
Contig10	SEQ ID N°3465
Contig11	SEQ ID N°3466
Contig12	SEQ ID N°3467
Contig13	SEQ ID N°3468
Contig14	SEQ ID N°3469
Contig15	SEQ ID N°3470
Contig16	SEQ ID N°3471
Contig17	SEQ ID N°3472
Contig18	SEQ ID N°3473
Contig19	SEQ ID N°3474
Contig20	SEQ ID N°3475
Contig21	SEQ ID N°3476
Contig22	SEQ ID N°3477
Contig23	SEQ ID N°3478
Contig24	SEQ ID N°3479
Contig25	SEQ ID N°3480
Contig26	SEQ ID N°3481
Contig27	SEQ ID N°3482
Contig28	SEQ ID N°3483
Contig29	SEQ ID N°3484
Contig30	SEQ ID N°3485
Contig31	SEQ ID N°3486
Contig32	SEQ ID N°3487
Contig33	SEQ ID N°3488
Contig34	SEQ ID N°3489
Contig35	SEQ ID N°3490
Contig36	SEQ ID N°3491
Contig37	SEQ ID N°3492
Contig38	SEQ ID N°3493
Contig39	SEQ ID N°3494
Contig40	SEQ ID N°3495
Contig41	SEQ ID N°3496
Contig42	SEQ ID N°3497
Contig43	SEQ ID N°3498
Contig44	SEQ ID N°3499

Contig45	SEQ ID N°3500
Contig46	SEQ ID N°3501
Contig47	SEQ ID N°3502
Contig48	SEQ ID N°3503
Contig49	SEQ ID N°3504
Contig50	SEQ ID N°3505
Contig51	SEQ ID N°3506

<u>Table V</u>: Surface proteins of *Legionella pneumophila* Paris strain

The proteins of surfaces specific to the Paris strain are indicated in bold.

SEQ ID	IPF	Annotation/simiarity to other proteins
3410	94.7	Surface protein, adhesion protein of <i>Streptococcus</i> sp. and <i>Pseudomonas</i> sp., Rtx toxin
704	202.3	Surface antigen of Bordetella sp. and Coxiella burnetii
746		Surface protein of Wolbachia sp.
2267	440.4	Surface protein of Mycoplasma hominis, Streptococcus
2751	514.5	Protein similar to 440.4
3192	627.1	Surface protein of Streptococcus pyogenes (« collagen-like »)
3218	663.2	Lipopolysaccharide biosynthesis, O-antigen acetylase <i>Pseudomonas</i> aeruginosa
3221	667.3	Surface antigen of Trypanosoma cruzi
3222	668.4	IcmE of Legionella pneumophila
3317	803.3	Flagellar protein (« L-ring protein »)
3324	817.7	Surface protein of Mycoplasma hominis
136	1115.4	Rtx toxin of Magnetococcus sp., putative lipoprotein of Leptospira kirschneri
171	1171.3	Protein of « surface exclusion » type of Salmonella typhimurium
310		Transporter of protons of Coxiella burnetii
337	1429.4	Rtx toxin, surface protein of Bacillus cereus
481	1653.3	Activator of plasminogene of <i>Yersinia pestis</i> , protease associated with the cellular envelope
527	1724.3	Hydrolase of the cellular envelope of <i>Pseudomonas putida</i>
652	1910.6	Protein similar to 440.4
664	1933.4	Protein similar to 440.4
893		Surface antigen of <i>Rickettsia</i> sp.
972	2448.3	Hypothetical protein of Coxiella burnetii, periplasmic protein
1148		Surface protein of Spirochète
1298		FimV, assembly of pili of Legionella pneumophila
1361		Immunogene protein of Legionella pneumophila
1503		O-antigen acetylase of Pseudomnas aeruginosa
1521		Agglutinine, adhesine, surface protein of Brucella melitensis
1576		Surface antigen of Magnetospirillum magnetotacticum
1651		Glycoprotein rich in histidine of <i>Plasmodium lophurae</i>
1755		Lipoprotein pal of Legionella pneumophila
1847	3780.2	Surface protein of Plasmodium falciparum

1877	3827.2	O-antigen acetylase of Pseudomnas aeruginosa
2224	4347.2	Protoin of the outernal mountains of D. 11 11 1 C
2224	4347.2	antigen of Rickettsia sp.
2406	4608.1	<u> </u>
2843	5349.3	Major surface protein of Anaplasma marginale,
2043		hypothetical protein of Plasmodium falciparum
2930	5526.2	Adhesine, virulence protein of Escherichia coli
3037 5739.		Protein of « surface exclusion » type of Pseudomonas putida,
3037	3739.3	Enterococcus faecalis, Rtx toxin
3139	6037.1	Surface antigen of Entamoeba histolytica and Plasmodium falciparum
3157	6079.1	Surface antigen of Trypanosoma cruzi
3165	6097.1	
3181	6131.1	Adhesing/surface mustain of D.
3101	0131.1	Enterococcus faecalis, Rtx toxin

<u>Table VI</u>: Proteins implied in biosynthesis of polysaccharides having a cellular envelope of *Legionella pneumophila*

SEQ I	D IPF	Annotation/similarity with other proteins
112	269.1	Hantagyl transferage his grantle size of live along 1 1 1 C + H
321	8 663.2	O-acetyl transferase, modification of lipopolysaccharides in <i>Vibrio cholerae</i>
28	1360.6	Protein implied in biosynthesis of lipopolysaccharides in Methanosarcina
63	2 1882.2	Polysaccharide deacetylase of Coxiella burnetii
91		Proteine CapM of Rickettsia conorii, glycosyltransferase
150		A cotal con of cution O CD 1
155	5 3348.2	Predicted xylanase/chitine deacetylase of Cytophaga hutchinsonii
187	7 3827.2	A potulage of outleast O of Day 1
192	8 3923.2	Detential animary of an in a side of the s
196	3 3980.1	Phosphopantetheine adenylyltransferase of <i>Ralstonia metallidurans</i> , biosynthesis of lipopolysaccharides
220	4 4323.1	
221	2 4334.1	Polysaccharide deacetylase, xylanase/chitin deacetylase
224	3 4371.1	Polysaccharide deacetylase, xylanase/chitin deacetylase
232	4 4488.1	Aminotransferase, synthesis of lipopolysaccharides
237	8 4567.2	WciT of Streptococcus pneumoniae, biosynthesis of polysaccharides
241		Biosynthesis of antigen O, hypothetical protein of Coxiella burnetii
241	1 4618.1	Biosynthesis of lipopolysaccharides, glycosyltransferase
5		

Table X: Correspondence of the numbers attributed to the contigs of L. pneumophila philadelphia with the numbers of the SEQ ID identified in the list of sequences

Contig1	SEQ ID N∞7061
Contig2	SEQ ID N∞7062
Contig3	SEQ ID N∞7063
Contig4	SEQ ID N∞7064
Contig5	SEQ ID N∞7065
Contig6	SEQ ID N∞7066
Contig7	SEQ ID N∞7067
Contig8	SEQ ID N∞7068
Contig9	SEQ ID N∞7069
Contig10	SEQ ID N∞7070
Contig11	SEQ ID N∞7071
Contig12	SEQ ID N∞7072
Contig13	SEQ ID N∞7073

Table XI: List of the sequences of L. pneumophila philadelphia identified as specific to 5 this strain relative to the Paris and Lens strains and position of these sequences on the contigs

Indication on the specifics of the Philadelphia strain

IPF Lp Philadelphia	Contig	SEQ ID	Position1	Position2
1007.1	CONTIG9	SEQ ID N°7069	1062411	1062962
10563.1	CONTIG13	SEQ ID N°7073	1463	2446
3775.3	CONTIG13	SEQ ID N°7073	1463	2446
1067.1	CONTIG7	SEQ ID N°7067	163133	163567
1980.3	CONTIG7	SEQ ID N°7067	163628	163918
1102.1	CONTIG7	SEQ ID N°7067	189792	190835
1109.1	CONTIG7	SEQ ID N°7067	195874	198036
4935.1	CONTIG9	SEQ ID N°7069	1604318	1605460
7686.1	CONTIG9	SEQ ID N°7069	1604318	1605460
1771.2	CONTIG8	SEQ ID N°7068	552424	553377
1773.1	CONTIG9	SEQ ID N°7069	961264	962745
1296.1	CONTIG9	SEQ ID N°7069	961264	962745
1297.1	CONTIG9	SEQ ID N°7069	959864	960817
1298.1	CONTIG9	SEQ ID N°7069	959562	959810
1302.1	CONTIG9	SEQ ID N°7069	958145	958699
1303.1	CONTIG9	SEQ ID N°7069	957452	957922
1307.1	CONTIG9	SEQ ID N°7069	956035	956523
1309.1	CONTIG9	SEQ ID N°7069	955209	955589
1310.1	CONTIG9	SEQ ID N°7069	954726	955034
1312.1	CONTIG9	SEQ ID N°7069	953857	954711
1313.1	CONTIG9	SEQ ID N°7069	953085	953864
1315.1	CONTIG9	SEQ ID N°7069	952598	953161

1319.1	CONTIG9	SEQ ID N°7069	950926	951858
1320.1	CONTIG9	SEQ ID N°7069	948772	950907
1321.1	CONTIG9	SEQ ID N°7069	948180	948743
1322.1	CONTIG9	SEQ ID N°7069	947726	948187
1323.1	CONTIG9	SEQ ID N°7069	947134	947706
1324.1	CONTIG9	SEQ ID N°7069	946011	946685
1325.1	CONTIG9	SEQ ID N°7069	945182	945862
1327.1	CONTIG9	SEQ ID N°7069	944125	945009
1328.1	CONTIG9	SEQ ID N°7069	943800	944288
1330.1	CONTIG9	SEQ ID N°7069	943233	943535
1331.1	CONTIG9	SEQ ID N°7069	942938	943243
1332.1	CONTIG9	SEQ ID N°7069	942371	942934
1333.1	CONTIG9	SEQ ID N°7069	941655	942368
1334.1	CONTIG9	SEQ ID N°7069	940367	941653
1337.1	CONTIG9	SEQ ID N°7069	940003	940242
1339.1	CONTIG9	SEQ ID N°7069	937439	940036
1340.1	CONTIG9	SEQ ID N°7069	937111	937446
1341.1	CONTIG9	SEQ ID N°7069	936506	937114
1342.1	CONTIG9	SEQ ID N°7069	935514	936509
1343.1	CONTIG9	SEQ ID N°7069	934830	935513
1344.1	CONTIG9	SEQ ID N°7069	933019	934890
1345.1	CONTIG9	SEQ ID N°7069	932244	933026
1346.1	CONTIG9	SEQ ID N°7069	931832	932254
1347.1	CONTIG9	SEQ ID N°7069	930450	931835
2328.1	CONTIG3	SEQ ID N°7063	749	1696
1348.1	CONTIG9	SEQ ID N°7069	927724	930444
1350.1	CONTIG9	SEQ ID N°7069	927256	927720
1353.1	CONTIG9	SEQ ID N°7069	925299	927188
1355.1	CONTIG9	SEQ ID N°7069	919333	925278
1356.1	CONTIG9	SEQ ID N°7069	918010	919137
1358.1	CONTIG9	SEQ ID N°7069	916581	917417
1359.1	CONTIG9	SEQ ID N°7069	916079	916570
1360.1	CONTIG9	SEQ ID N°7069	914469	916049
1361.1	CONTIG9	SEQ ID N°7069	913897	914271
1365.1	CONTIG9	SEQ ID N°7069	911529	913049
1366.1	CONTIG9	SEQ ID N°7069	910007	911458
1367.1	CONTIG9	SEQ ID N°7069	909405	909701
1368.1	CONTIG9	SEQ ID N°7069	908527	909309
1369.1	CONTIG9	SEQ ID N°7069	908004	908504
1370.1	CONTIG9	SEQ ID N°7069	907151	907993
1371.1	CONTIG9	SEQ ID N°7069	905264	907069
1373.1	CONTIG9	SEQ ID N°7069	903498	905264
1376.1	CONTIG9	SEQ ID N°7069	902217	902987
1378.1	CONTIG9	SEQ ID N°7069	900968	901987
1380.1	CONTIG9	SEQ ID N°7069	899639	900985
1382.1	CONTIG9	SEQ ID N°7069	898691	899314
1383.1	CONTIG9	SEQ ID N°7069	898266	898532
1384.1	CONTIG9	SEQ ID N°7069	897979	898281
1386.1	CONTIG9	SEQ ID N°7069	897062	898066
1435.1	CONTIG9	SEQ ID N°7069	864915	866060
		•		

1492.1	CONTIG9	SEQ ID N°7069	821283	821579
1494.1	CONTIG9	SEQ ID N°7069	820792	821109
1500.1	CONTIG9	SEQ ID N°7069	814942	815592
1501.1	CONTIG9	SEQ ID N°7069	814765	815241
1503.1	CONTIG9	SEQ ID N°7069	813201	813485
1518.1	CONTIG9	SEQ ID N°7069	800919	803852
1519.1	CONTIG9	SEQ ID N°7069	799375	800655
1520.1	CONTIG9	SEQ ID N°7069	796924	799023
1538.1	CONTIG9	SEQ ID N°7069	775795	776787
1627.1	CONTIG8	SEQ ID N°7068	454194	455027
1631.1	CONTIG8	SEQ ID N°7068	456917	457408
1635.1	CONTIG8	SEQ ID N°7068	459876	460751
1663.1	CONTIG8	SEQ ID N°7068	477274	477810
1674.1	CONTIG8	SEQ ID N°7068	488306	490213
1676.1	CONTIG8	SEQ ID N°7068	491037	491288
1723.1	CONTIG8	SEQ ID N°7068	522733	523200
1724.1	CONTIG8	SEQ ID N°7068	523306	523560
1725.1	CONTIG8	SEQ ID N°7068	523670	523945
1749.1	CONTIG8	SEQ ID N°7068	537066	537341
1760.1	CONTIG8	SEQ ID N°7068	544546	545574
1767.1	CONTIG8	SEQ ID N°7068	549553	550482
1772.1	CONTIG6	SEQ ID N°7066	779	1804
1785.1	CONTIG9	SEQ ID N°7069	970464	970808
1787.1	CONTIG9	SEQ ID N°7069	970976	971416
1892.1	CONTIG7	SEQ ID N°7067	76315	77967
3771.3	CONTIG13	SEQ ID N°7073	43581	46268
1952.1	CONTIG9	SEQ ID N°7069	1328145	1328393
1981.1	CONTIG9	SEQ ID N°7069	1347920	1348357
2005.1	CONTIG9	SEQ ID N°7069	1364695	1366614
2006.1	CONTIG9	SEQ ID N°7069	1366816	1368231
2026.1	CONTIG9	SEQ ID N°7069	1380350	1380739
2059.1	CONTIG9	SEQ ID N°7069	1407671	1408075
2066.1	CONTIG9	SEQ ID N°7069	1418764	1421481
2083.1	CONTIG9	SEQ ID N°7069	1436828	1439185
2084.1	CONTIG9	SEQ ID N°7069	1439377	1440240
2086.1	CONTIG9	SEQ ID N°7069	1440346	1441680
2125.2	CONTIG9	SEQ ID N°7069	1470612	1471472
2132.1	CONTIG11	SEQ ID N°7071	144883	145770
2133.1	CONTIG11	SEQ ID N°7071	144142	144882
2134.1	CONTIG11	SEQ ID N°7071	143221	143718
2135.1	CONTIG11	SEQ ID N°7071	143204	144145
2141.1	CONTIG11	SEQ ID N°7071	137310	137879
2202.1	CONTIG11	SEQ ID N°7071	87995	89587
2311.1	CONTIG8	SEQ ID N°7068	553457	554560
2312.1	CONTIG8	SEQ ID N°7068	554678	555862
2314.1	CONTIG8	SEQ ID N°7068	555930	556454
2315.1	CONTIG8	SEQ ID N°7068	556495	556839
2316.1	CONTIG8	SEQ ID N°7068	556651	558360
2317.1	CONTIG8	SEQ ID N°7068	558425	558796
2319.1	CONTIG8	SEQ ID N°7068	559260	560681

2321.1	CONTIG8	SEQ ID N°7068	560851	562347
2323.1	CONTIG8	SEQ ID N°7068	562541	562981
2324.1	CONTIG8	SEQ ID N°7068	563118	563813
2325.1	CONTIG8	SEQ ID N°7068	563899	564930
2326.1	CONTIG8	SEQ ID N°7068	564927	566123
2330.1	CONTIG8	SEQ ID N°7068	567385	567954
2333.1	CONTIG8	SEQ ID N°7068	568120	569727
2337.1	CONTIG8	SEQ ID N°7068	572901	573461
2343.1	CONTIG8	SEQ ID N°7068	575955	578648
2345.1	CONTIG8	SEQ ID N°7068	578635	579891
2346.1	CONTIG8	SEQ ID N°7068	579888	581024
2348.1	CONTIG8	SEQ ID N°7068	581063	584191
2350.1	CONTIG8	SEQ ID N°7068	584584	586371
2357.1	CONTIG8	SEQ ID N°7068	590170	591033
2373.1	CONTIG8	SEQ ID N°7068	601883	602209
2385.1	CONTIG8	SEQ ID N°7068	613763	615529
2465:1	CONTIG12	SEQ ID N°7072	96957	97913
2466.1	CONTIG12	SEQ ID N°7072	97888	98478
2467.1	CONTIG12	SEQ ID N°7072	98499	99803
2468.1	CONTIG12	SEQ ID N°7072	99820	100638
2469.1	CONTIG12	SEQ ID N°7072	100647	101573
2470.1	CONTIG12	SEQ ID N°7072	101679	102866
2471.1	CONTIG12	SEQ ID N°7072	103108	104379
2472.1	CONTIG12	SEQ ID N°7072	104380	105558
2473.1	CONTIG12	SEQ ID N°7072	105588	106370
2498.1	CONTIG12	SEQ ID N°7072	124135	124545
2500.1	CONTIG12	SEQ ID N°7072	123924	126110
2684.2	CONTIG8	SEQ ID N°7068	740173	740430
2710.1	CONTIG9	SEQ ID N°7069	1778548	1780281
2743.1	CONTIG9	SEQ ID N°7069	1810120	1810353
2769.1	CONTIG9	SEQ ID N°7069	1826652	1828562
2782.3	CONTIG8	SEQ ID N°7068	78469	79875
5631.2	CONTIG8	SEQ ID N°7068	78469	79875
2784.1	CONTIG8	SEQ ID N°7068	73057	73398
2889.1	CONTIG13	SEQ ID N°7073	19409	20020
2890.1	CONTIG13	SEQ ID N°7073	18305	19162
2892.1	CONTIG13	SEQ ID N°7073	17653	17907
2894.1	CONTIG13	SEQ ID N°7073	15676	17652
2895.1	CONTIG13	SEQ ID N°7073	15195	15683
2896.1	CONTIG13	SEQ ID N°7073	14143	14952
3353.1	CONTIG13	SEQ ID N°7073	14143	14952
2915.1	CONTIG13	SEQ ID N°7073	21609	22910
3909.2	CONTIG13	SEQ ID N°7073	21609	22910
2916.1	CONTIG13	SEQ ID N°7073	23001	24515
3908.2	CONTIG13	SEQ ID N°7073	23001	24515
2917.1	CONTIG13	SEQ ID N°7073	24645	24908
2918.1	CONTIG13	SEQ ID N°7073	24971	26290
2919.1	CONTIG13	SEQ ID N°7073	26453	26965
2920.1	CONTIG13	SEQ ID N°7073	27050	27442
2921.1	CONTIG13	SEQ ID N°7073	27535	28215
		`	*	

3018.1	CONTIG13	SEQ ID N°7073	41565	42278
10565.1	CONTIG9	SEQ ID N°7069	1320956	1322047
1947.2	CONTIG9	SEQ ID N°7069	1320956	1322047
3019.1	CONTIG9	SEQ ID N°7069	1320956	1322047
3020.1	CONTIG9	SEQ ID N°7069	1320447	1320905
3021.1	CONTIG9	SEQ ID N°7069	1319365	1320159
3023.1	CONTIG9	SEQ ID N°7069	1317520	1319235
3024.1	CONTIG9	SEQ ID N°7069	1316653	1317507
3025.1	CONTIG9	SEQ ID N°7069	1315445	1316551
3027.1	CONTIG9	SEQ ID N°7069	1314906	1315190
3046.1	CONTIG9	SEQ ID N°7069	1298926	1299282
3047.1	CONTIG9	SEQ ID N°7069	1298615	1299712
3087.1	CONTIG9	SEQ ID N°7069	1016602	1017420
3097.1	CONTIG11	SEQ ID N°7071	183622	184182
3100.1	CONTIG11	SEQ ID N°7071	182447	182740
3101.1	CONTIG11	SEQ ID N°7071	182080	182409
3102.1	CONTIG11	SEQ ID N°7071	180595	181965
3103.1	CONTIG11	SEQ ID N°7071	179123	180463
3104.1	CONTIG11	SEQ ID N°7071	177735	179093
3105.1	CONTIG11	SEQ ID N°7071	176469	177605
3317.1	CONTIG9	SEQ ID N°7069	1863795	1864346
2685.2	CONTIG13	SEQ ID N°7073	12622	13683
3356.1	CONTIG13	SEQ ID N°7073	12622	13683
3357.1	CONTIG13	SEQ ID N°7073	11421	12638
3362.1	CONTIG13	SEQ ID N°7073	7035	7835
3364.1	CONTIG13	SEQ ID N°7073	5298	6497
3365.1	CONTIG13	SEQ ID N°7073	4728	5141
34.1	CONTIG9	SEQ ID N°7069	40333	43707
35.1	CONTIG9	SEQ ID N°7069	44063	44371
3538.1	CONTIG9	SEQ ID N°7069	1883947	1885137
3539.1	CONTIG9	SEQ ID N°7069	1885249	1885500
3541.1	CONTIG9	SEQ ID N°7069	1887373	1888815
3542.1	CONTIG9	SEQ ID N°7069	1888997	1889311
3544.1	CONTIG9	SEQ ID N°7069	1889663	1889962
3545.1	CONTIG9	SEQ ID N°7069	1890063	1890284
3548.1	CONTIG9	SEQ ID N°7069	1891102	1892148
3550.2	CONTIG9	SEQ ID N°7069	1893944	1895122
3551.1	CONTIG9	SEQ ID N°7069	1895356	1895775
3552.2	CONTIG9	SEQ ID N°7069	1895768	1896094
3683.1	CONTIG8	SEQ ID N°7068	129540	131426
37.1	CONTIG9	SEQ ID N°7069	44561	44827
3740.2	CONTIG9	SEQ ID N°7069	1025799	1027007
2327.1	CONTIG12	SEQ ID N°7072	10	987
3742.1	CONTIG12	SEQ ID N°7072	10	987
1944.1	CONTIG13	SEQ ID N°7073	999	1205
3793.1	CONTIG9	SEQ ID N°7069	1491228	1491776
3794.1	CONTIG9	SEQ ID N°7069	1490472	1491212
3795.4	CONTIG9	SEQ ID N°7069	1489860	1490459
3797.2	CONTIG9	SEQ ID N°7069	1921869	1923518
3798.1	CONTIG9	SEQ ID N°7069	1923682	1924194

3799.1	CONTIG9	SEQ ID N°7069	1924210	1924524
3800.1	CONTIG9	SEQ ID N°7069	1924550	1926376
3803.1	CONTIG9	SEQ ID N°7069	1926518	1927483
3827.1	CONTIG13	SEQ ID N°7073	2622	3332
3829.2	CONTIG13	SEQ ID N°7073	3307	4557
3847.1	CONTIG9	SEQ ID N°7069	1482616	1483233
3848.1	CONTIG9	SEQ ID N°7069	1481860	1482336
3849.3	CONTIG9	SEQ ID N°7069	1480223	1481680
3850.1	CONTIG9	SEQ ID N°7069	1479788	1480216
3890.1	CONTIG13	SEQ ID N°7073	20841	21641
39.1	CONTIG9	SEQ ID N°7069	44906	45655
8044.1	CONTIG7	SEQ ID N°7067	16618	17850
3986.2	CONTIG5	SEQ ID N°7065	2	2959
4124.2	CONTIG5	SEQ ID N°7065	2	2959
3997.1	CONTIG9	SEQ ID N°7069	1920982	1921893
40.1	CONTIG9	SEQ ID N°7069	45836	46858
4018.1	CONTIG9	SEQ ID N°7069	1913464	1914630
4019.1	CONTIG9	SEQ ID N°7069	1912844	1913476
4022.2	CONTIG9	SEQ ID N°7069	1911062	1911376
4064.1	CONTIG9	SEQ ID N°7069	1486212	1486631
4065.1	CONTIG9	SEQ ID N°7069	1485967	1486812
4066.1	CONTIG9	SEQ ID N°7069	1486860	1487468
4067.4	CONTIG9	SEQ ID N°7069	1487626	1488717
41.1	CONTIG9	SEQ ID N°7069	46950	47837
43.1	CONTIG9	SEQ ID N°7069	48387	50369
4936.1	CONTIG9	SEQ ID N°7069	1605170	1605475
504.1	CONTIG9	SEQ ID N°7069	473249	474757
506 .1	CONTIG9	SEQ ID N°7069	475061	476458
507.1	CONTIG9	SEQ ID N°7069	476469	477350
508.1	CONTIG9	SEQ ID N°7069	477545	477874
518.1	CONTIG9	SEQ ID N°7069	485931	487241
519.1	CONTIG9	SEQ ID N°7069	487186	487464
521.1	CONTIG9	SEQ ID N°7069	487597	487989
669.1	CONTIG9	SEQ ID N°7069	604354	605595
6736.1	CONTIG6	SEQ ID N°7066	1824	2537
6874.1	CONTIG8	SEQ ID N°7068	575999	576325
750.1	CONTIG9	SEQ ID N°7069	1271952	1274519
751.1	CONTIG9	SEQ ID N°7069	1271377	1271823
752.1	CONTIG9	SEQ ID N°7069	1270905	1271255
760.1	CONTIG9	SEQ ID N°7069	1266574	1267239
761.1	CONTIG9	SEQ ID N°7069	1266303	1266707
764.1	CONTIG9	SEQ ID N°7069	1263381	1263866
765.1	CONTIG9	SEQ ID N°7069	1261771	1262709
774.1	CONTIG9	SEQ ID N°7069	1258317	1260068
778.1	CONTIG9	SEQ ID N°7069	1256902	1257267
8067.2	CONTIG9	SEQ ID N°7069	1488930	1489379
8073.2	CONTIG9	SEQ ID N°7069	1484592	1484990
10294.1	CONTIG8	SEQ ID N°7068	119556	120962
7817.1	CONTIG8	SEQ ID N°7068	119556	120962
8134.1	CONTIG8	SEQ ID N°7068	119556	120962

874.1	CONTIG9	SEQ ID N°7069	1177144	1178064
875.1	CONTIG9	SEQ ID N°7069	1176375	1176995
876.1	CONTIG9	SEQ ID N°7069	1175483	1176361
1769.1	CONTIG3	SEQ ID N°7063	21	716
9388.1	CONTIG3	SEQ ID N°7063	21	716
4934.1	CONTIG9	SEQ ID N°7069	1819527	1820453
2798.2	CONTIG8	SEQ ID N°7068	71193	77576
3630.3	CONTIG12	SEQ ID N°7072	11054	16705

<u>Table XII</u>: Position of former contigs of the Paris strain on the genomic sequence of the chromosome of the Paris strain of sequence SEQ ID 3507 and of the plasmid of the Paris strain of sequence SEQ ID 3508

	pos1	pos2	former contig	SEQ ID of former contig
chromosome	1	44600	41	SEQ ID N∞41
	44600	161000	56	SEQ ID N∞56
	162000	223000	54	SEQ ID N∞54
	223000	232000	39	SEQ ID N∞39
•	236000	424000	49	SEQ ID N∞49
	437000	469000	37	SEQ ID N∞37
	464000	758000	53	SEQ ID N∞53
	758000	781000	45	SEQ ID N∞45
	789000	879000	43	SEQ ID N∞43
-	883000	901000	36	SEQ ID N∞36
	898000	986000	42	SEQ ID N∞42
	990000	1160000	.46	SEQ ID N ∞46
	1160000	1214000	39	SEQ ID N∞39
	1214000	1352000	54	SEQ ID N∞54
	1352000	1670000	52	SEQ ID N∞52
	1670000	1736000	40	SEQ ID N∞40
	1736000	2040000	55	SEQ ID N∞55
	2044000	2093000	56	SEQ ID N∞56
	2093000	2204722	45	SEQ ID N∞45
	2204000	2298000	50	SEQ ID N∞50
	2298000	2656000	56	SEQ ID N∞56
	2656000	2740000	50	SEQ ID N∞50
•	2740000	2753600	33	SEQ ID N∞33
	2753600	2954000	47	SEQ ID N∞47
	2954000	3178000	51	SEQ ID N∞51
	3178000	3289000	44	SEQ ID N∞44
	3289000	3449000	48	SEQ ID N∞48
	3449000	3463000	34	SEQ ID N∞34
	3463000	3503610	41	SEQ ID N∞41
plasmid	1	131900	55(position1 to 132400)	SEQ ID N∞55

<u>Table XIII</u>: Correspondence of the numbers attributed to the chromosome and au plasmid of the Paris and Lens strain with the SEQ ID numbers identified in the list of sequences

5	SeqID=3507	chromosome of the Paris strain
	SeqID=3508	plasmid of the Paris strain
•	SeqID=6733	chromosome of the Lens strain
	SeqID=6734	plasmid of the Lens strain

Table XIV: Correspondence of the numbers attributed to the genes of the Paris strain on its chromosome of sequence SEQ ID 3507 and on its plasmid of sequence SEQ ID 3508 with the numbers of the SEQ ID identified in the list of sequences and position of the nucleic sequences coding these genes on the sequence of the chromosome and of the plasmid with their putative function

15

Table XV: Nature of the class listed in the "Class" column in Tables XIV and XVI

- 1. Cellular envelope and cellular processes
- 20 1.1 Cellular wall and external membrane
 - 1.2 Proteins of transport/bond and lipoproteins
 - 1.3 Sensors (transduction of signal)
 - 1.4 Bioenergy of membrane
 - 1.5 Mobility and chimiotaxia
- 25 1.6 Secretion of protein
 - 1.7 Cellular division
 - 1.8 Structures of cellular surface and pili
 - 2. Intermediary metabolism

30

- 2.1 Metabolism of glucides and related molecules
- 2.1.1 Specific ways
- 2.1.2 Principal glycolytic ways
- 2.1.3 TCA cycle
- 2.2 Metabolism of aminoacids and related molecules
 - 2.3 Metabolism of nucleotides and nucleic acids
 - 2.4 Metabolism of lipids
 - 2.5 Metabolism of coenzymes and prosthetic groups
 - 2.6 Metabolism of phosphate

40

- 3. Information paths
- 3.1 Replication of DNA
- 3.2 Repair and restriction/modification of DNA

	3.3 3.4 3.5	Recombination of DNA Segregation and encapsidation of DNA Synthesis of RNA
5	3.5.1 3.5.2 3.5.3 3.5.4 3.6	Initiation Regulation Elongation Termination Modification of RNA
10	3.7 3.7.1 3.7.2 3.7.3 3.7.4	Synthesis of protein Ribosomal proteins Synthetases of aminoacyl-tRNA Initiation
15	3.7.5 3.8 3.9	Elongation Termination Modification of protein Folding of protein
	4.	Other functions
20 25	4.1 4.2 4.3 4.4 4.5 4.6	Adaptation to atypical conditions Detoxification Toxins Functions relating to phage Transposon, IS, Plasmid Various
	5.	Similar to unknown proteins
30	5.1 5.2	Of Legionella (similar though not the same) Of other organisms
	6.	No similarity
35	Similar Similar Similar	r to the enzyme IIN of the PTS XXXX-specific system r to the transcriptional regulator (family xx) r to the transportor ABC (protein of ATP bond) r to the transportor ABC (permease) r to the transportor ABC (bond protein)
40	Similar Similar	r to the response regulator with two compounds r to the histidine kinase sensor with two compounds n bound to putative peptidoglycane (LPXTG pattern)

Table XVI: Correspondence of the numbers attributed to the specific genes of the Lens strain relative to the Paris and Philadelphia strains on its chromosome of sequence SEQ ID 6733 and on its plasmid of sequence SEQ ID 6734 with the numbers of SEQ ID identified in the list of sequences and position of the nucleic sequences coding these genes on the sequence of the chromosome and the plasmid with their putative function

<u>Table XVII</u>: List of the specific sequences of the Paris strain relative to the Lens and Philadelphia strains with their Pasteur Institute « ORF » correspondence number and accession number in the gene banks

```
1056.1 SEQID=3544 EMBL NAME=lpp1800
1067.1 SEQID=3551 EMBL NAME=lpp1877
1069.2 SEQID=3552 EMBL NAME=lpp1878
1076.3 SEQID=6591 EMBL NAME=plpp0105
1077.1 SEQID=6592 EMBL NAME=plpp0104
1078.2 SEQID=6593 EMBL NAME=plpp0103
1080.2 SEQID=3558 EMBL NAME=lpp3012
1081.2 SEQID=3559 EMBL NAME=lpp3011
11.2
      SEQID=3573 EMBL NAME=lpp0196
114.2
      SEQID=3598 EMBL NAME=lpp2957
115.2
      SEQID=3603 EMBL NAME=lpp2956
      SEQID=3609 EMBL NAME=lpp2955
116.1
1160.3 SEQID=3610 EMBL NAME=lpp0125
1171.4 SEQID=6594 EMBL NAME=plpp0017
1172.2 SEQID=6595 EMBL NAME=plpp0018
      SEQID=3618 EMBL NAME=lpp2954
118.1
1183.4 SEQID=3621 EMBL NAME=lpp0356
1213.2 SEQID=3638 EMBL NAME=lpp0257
1235.3 SEQID=6598 EMBL NAME=plpp0121
1237.2 SEQID=6600 EMBL NAME=plpp0119
1299.3 SEQID=6601 EMBL NAME=plpp0036
13.1
      SEQID=3688 EMBL NAME=lpp0195
1342.3 SEQID=6602 EMBL NAME=plpp0034
1344.4 SEQID=6603 EMBL NAME=plpp0033
1362.3 SEQID=6604 EMBL NAME=plpp0098
1364.2 SEQID=6605 EMBL NAME=plpp0099
1372.2 SEQID=3726 EMBL NAME=lpp2385
1373.1 SEQID=3727 EMBL NAME=lpp2384
1375.2 SEQID=3728 EMBL NAME=lpp2383
1376.2 SEQID=3729 EMBL NAME=lpp2382
1387.2 SEQID=3735 EMBL NAME=lpp0079
1388.2 SEQID=3736 EMBL NAME=lpp0080
139.6
      SEQID=3737 EMBL NAME=lpp1100
1392.2 SEQID=3740 EMBL NAME=lpp1097
1394.2 SEQID=3741 EMBL NAME=lpp2557
1429.4 SEQID=3766 EMBL NAME=lpp2442
1522.2 SEQID=6606 EMBL NAME=plpp0127
1523.2 SEQID=6607 EMBL NAME=plpp0128
1524.3 SEQID=6608 EMBL NAME=plpp0129
1566.3 SEQID=3846 EMBL NAME=lpp2490
1570.4 SEQID=3850 EMBL NAME=lpp0077
1599.5 SEQID=6610 EMBL NAME=plpp0039
1623.4 SEQID=3881 EMBL NAME=lpp2394
1624.5 SEQID=3882 EMBL NAME=lpp2395
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SEQID=3887 EMBL_NAME=lpp2344
  163.1
  1655.2 SEQID=3905 EMBL_NAME=lpp1895
  1683.2 SEQID=3923 EMBL_NAME=lpp0046
       SEQID=3946 EMBL_NAME=lpp2978
  172.1
 1735.1 SEQID=3955 EMBL_NAME=lpp1862
 1761.4 SEQID=6611 EMBL_NAME=plpp0074
 1779.3 SEQID=3985 EMBL_NAME=lpp2040
 1787.4 SEQID=3992 EMBL_NAME=lpp1824
       SEQID=4000 EMBL_NAME=lpp0192
 18.1
 1803.2 SEQID=4003 EMBL_NAME=lpp2374
 1815.2 SEQID=4010 EMBL_NAME=lpp2986
 1848.5 SEQID=6613 EMBL_NAME=plpp0124
 1849.4 SEQID=6614 EMBL_NAME=plpp0125
 1852.2 SEQID=6615 EMBL_NAME=plpp0126
 1891.2 SEQID=4056 EMBL_NAME=lpp3007
 19.1
       SEQID=4061 EMBL_NAME=lpp0191
 1920.3 SEQID=4074 EMBL_NAME=lpp2405
 1923.2 SEQID=4075 EMBL NAME=lpp2406
 1924.4 SEQID=4076 EMBL_NAME=lpp2407
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<u>Table XVIII</u>: List of the specific sequences of the Lens strain relative to the Paris and Philadelphia strains with their «ORF» Institut Pasteur correspondence number and accession number in the gene banks

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1237.2	SEQID=6770	EMBL_NAME=1p12867
1251.1	SEQID=6773	EMBL_NAME=1pl2857
1252.1	SEQID=6774	EMBL_NAME=1p12848
1255.2	SEQID=6775	EMBL_NAME=1p12847
1258.1	SEQID=6776	EMBL_NAME=1p12845
126.1		EMBL_NAME=lpl2843
120.1	SEQID=6777	EMBL_NAME=lpl0165
127.1	SEQID=6778	EMBL_NAME=lpl0164
1274.1	SEQID=6779	EMBL_NAME=1p12842
1275.1	SEQID=6780	EMBL_NAME=lpl2841
1278.1	SEQID=6781	EMBL_NAME=1p12840
1278.1	SEQID=6782	EMBL_NAME=lpl2839
1279.1	SEQID=6783	EMBL_NAME=lpl2838
	SEQID=6784	EMBL_NAME=lpl2837
1283.1	SEQID=6786	EMBL_NAME=lpl2835
1284.1	SEQID=6787	EMBL_NAME=lpl2834
1285.1	SEQID=6788	EMBL_NAME=lpl2833
1296.1	SEQID=6789	EMBL_NAME=lpl2827
1297.1	SEQID=6790	EMBL_NAME=lp12826
1321.1	SEQID=6791	EMBL_NAME=lpl2806
1422.1	SEQID=6792	EMBL_NAME=lpl2741
1535.1	SEQID=6795	EMBL_NAME=lp10612
154.1	SEQID=6796	EMBL_NAME=lpl0146
156.1	SEQID=6797	EMBL_NAME=lpl0145
157.1	SEQID=6798	EMBL_NAME=lpl0144
159.1	SEQID=6799	EMBL_NAME=lpl0143
1697.1	SEQID=6800	EMBL_NAME=lpl0718
1718.1	SEQID=6801	EMBL_NAME=lpl0729
1824.1	SEQID=6803	EMBL_NAME=lpl0801
1826.1	SEQID=6805	EMBL_NAME=lp10803
1827.1	SEQID=6806	EMBL_NAME=lpl0804
1834.1	SEQID=6809	EMBL_NAME=lpl0811
1955.1	SEQID=6810	EMBL_NAME=lpl0904
2110.1	SEQID=6812	EMBL_NAME=lpl1015
2137.1	SEQID=6816	EMBL_NAME=lpl1037
2141.1	SEQID=6818	EMBL_NAME=lpl1041

2142.2	SEQID=6819	EMBL_NAME=lpl1042
2145.1	SEQID=6821	EMBL_NAME=lpl1043a
2152.2	SEQID=6823	EMBL_NAME=lpl1048
2155.1	SEQID=6824	EMBL_NAME=lpl1050
2170.1	SEQID=6827	EMBL_NAME=lpl1059
2180.1	SEQID=6828	EMBL NAME=lpl1067
2181.1	SEQID=6829	EMBL_NAME=lpl1068
2182.1	SEQID=6830	EMBL_NAME=lpl1069
2184.1	SEQID=6831	EMBL NAME=lpl1070
2186.2	SEQID=6832	EMBL NAME=lpl1071
2189.1	SEQID=6833	EMBL_NAME=lpl1073
2194.1	SEQID=6835	EMBL_NAME=lpl1076
2195.1	SEQID=6836	EMBL_NAME=lpl1077
2198.1	SEQID=6837	EMBL_NAME=lpl1080
2199.1	SEQID=6838	EMBL_NAME=lpl1081
2200.1	SEQID=6839	EMBL NAME=lpl1082
2201.1	SEQID=6840	EMBL NAME=lpl1083
2202.1	SEQID=6841	EMBL NAME=lpl1084
2203.1	SEQID=6842	EMBL NAME=lpl1085
2206.1	SEQID=6844	EMBL_NAME=lpl1087
2207.1	SEQID=6845	EMBL_NAME=lpl1088
2208.1	SEQID=6846	EMBL_NAME=lpl1089
2210.1	SEQID=6848	EMBL_NAME=lpl1091
2211.1	SEQID=6849	EMBL_NAME=lpl1092
2213.1	SEQID=6850	EMBL_NAME=lpl1093
2224.1	SEQID=6851	EMBL_NAME=lpl1101
2241.1	SEQID=6853	EMBL_NAME=lpl0199
2242.1	SEQID=6854	EMBL NAME=lpl0200
2245.1	SEQID=6856	EMBL NAME=lpl0202
2253.1	SEQID=6860	EMBL_NAME=lpl0207
2259.1	SEQID=6861	EMBL_NAME=lpl0211
2260.1	SEQID=6862	EMBL_NAME=lpl0212
2440.1	SEQID=6866	EMBL NAME=lpl2546
2441.1	SEQID=6867	EMBL_NAME=lpl2545
2516.1	SEQID=6868	EMBL_NAME=lpl2497
2518.1	SEQID=6870	EMBL_NAME=lpl2495
2521.1	SEQID=6871	EMBL_NAME=lpl2494
2523.1	SEQID=6872	EMBL_NAME=lpl2493
2524.1	SEQID=6873	EMBL_NAME=lpl2492
2525.1	SEQID=6874	EMBL_NAME=lpl2491
2526.1	SEQID=6875	EMBL_NAME=lpl2490
2527.1	SEQID=6876	EMBL_NAME=lpl2489
2529.1	SEQID=6877	EMBL_NAME=lpl2488
2530.1	SEQID=6878	EMBL_NAME=lp12487
2531.1	SEQID=6879	EMBL_NAME=lpl2486
2532.1	SEQID=6880	EMBL_NAME=lpl2485
2533.1	SEQID=6881	EMBL_NAME=lpl2484
2534.1	SEQID=6882	EMBL_NAME=lpl2483
2540.1	SEQID=6884	EMBL_NAME=lpl2477
2541.1	SEQID=6885	EMBL_NAME=lpl2476
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2547.1	SEQID=6886	EMBL_NAME=lpl2472
2584.1	SEQID=6888	EMBL_NAME=lpl2445
2640.1	SEQID=6890	EMBL_NAME=lpl2399
2658.1	SEQID=6891	EMBL_NAME=lpl2385
266.1	SEQID=6892	EMBL_NAME=lpl0069
267.1	SEQID=6894	EMBL_NAME=lpl0068
269.1	SEQID=6895	EMBL_NAME=lpl0067
270.1	SEQID=6897	EMBL_NAME=lpl0066
2701.1	SEQID=6898	EMBL_NAME=lp12354
2708.1	SEQID=6900	EMBL_NAME=lpl2350
2717.1	SEQID=6901	EMBL_NAME=lpl2344
2719.1	SEQID=6902	EMBL_NAME=lp12343
272.1	SEQID=6903	EMBL_NAME=lp10065
2720.1	SEQID=6904	EMBL_NAME=lpl0003
2722.1	SEQID=6905	EMBL_NAME=lpl2341
2723.1	SEQID=6906	
273.2	SEQID=6907	EMBL_NAME=lp12340
2738.1	SEQID=6908	EMBL_NAME=lpl1893
2749.1	SEQID=6909	EMBL_NAME=1pl2330
2775.1	SEQID=6910	EMBL_NAME=1p12323
2782.1	SEQID=6910	EMBL_NAME=1p12309
2795.1	SEQID=6913	EMBL_NAME=1p12305
2796.1	SEQID=6913	EMBL_NAME=1p12295
2798.1	SEQID=6915	EMBL_NAME=1pl2294
2801.1	SEQID=6916	EMBL_NAME=1p12293
2802.1	SEQID=6917	EMBL_NAME=1pl2292
2805.1	SEQID=6917 SEQID=6918	EMBL_NAME=1p12291
2806.1	SEQID=6918 SEQID=6919	EMBL_NAME=lp12289
2808.1	SEQID=6919 SEQID=6921	EMBL_NAME=1p12288
2810.1	SEQID=6923	EMBL_NAME=lpl2286
3002.1	_	EMBL_NAME=lpl2284
3042.1	SEQID=6925	EMBL_NAME=lpl2148
3054.1	SEQID=6926 SEQID=6927	EMBL_NAME=lpi2114
3056.1	SEQID=6927 SEQID=6928	EMBL_NAME=lp12107
3057.1	SEQID=6928 SEQID=6929	EMBL_NAME=lpl2106
3064.1	SEQID=6929 SEQID=6930	EMBL_NAME=lpl2105
3065.1	SEQID=6930 SEQID=6931	EMBL_NAME=lp12100
3066.1	SEQID=6931 SEQID=6932	EMBL_NAME=lpl2099
3140.1	_	EMBL_NAME=lpl2098
3155.1	SEQID=6933	EMBL_NAME=lpl2049
3156.1	SEQID=6935	EMBL_NAME=lpl2038
3158.1	SEQID=6936	EMBL_NAME=lpl2037 EMBL_NAME=lpl2036
3160.1	SEQID=6937	EMBL_NAME=lpl2036
3161.1	SEQID=6938	EMBL_NAME=lpl2035
3162.1	SEQID=6939	EMBL_NAME=lpl2034
3417.1	SEQID=6940	EMBL_NAME=lpl2033
3417.1 3420.1	SEQID=6942	EMBL_NAME=lpl0216
420.1 422.1	SEQID=6943	EMBL_NAME=lpl0217
	SEQID=6944	EMBL_NAME=lpl0218
435.1 728.1	SEQID=6945	EMBL_NAME=lpl0226
120.1	SEQID=6948	EMBL_NAME=lpl0552
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3743.2	SEQID=6949	EMBL_NAME=lp10565
3744.1	SEQID=6950	EMBL_NAME=lpl0566
3745.1	SEQID=6951	EMBL_NAME=lpl0567
3747.2	SEQID=6952	EMBL_NAME=lpl0568a
3748.2	SEQID=6953	EMBL_NAME=lpl0568b
3815.2	SEQID=6957	EMBL_NAME=plp10037
3818.1	SEQID=6959	EMBL_NAME=plp10039
3820.1	SEQID=6960	EMBL_NAME=plp10040
3822.1	SEQID=6961	EMBL_NAME=plp10042
3823.1	SEQID=6962	EMBL NAME=plp10043
3843.1	SEQID=6972	EMBL NAME=plpl0053
3848.2	SEQID=6973	EMBL_NAME=plpl0001
3850.1	SEQID=6974	EMBL_NAME=plpl0002
3863.1	SEQID=6978	EMBL_NAME=plpl0014
3865.1	SEQID=6979	EMBL_NAME=plpl0015
3866.1	SEQID=6980	EMBL_NAME=plpl0016
3868.1	SEQID=6981	EMBL_NAME=plpl0017
3870.1	SEQID=6982	EMBL_NAME=plpl0018
3871.1	SEQID=6983	EMBL_NAME=plpl0019
3873.1	SEQID=6984	EMBL_NAME=plpl0020
3874.1	SEQID=6985	EMBL NAME=plpl0021
3875.1	SEQID=6986	EMBL NAME=plp10022
3877.1	SEQID=6987	EMBL_NAME=plp10023
3878.1	SEQID=6988	EMBL_NAME=plpl0024
3880.1	SEQID=6989	EMBL_NAME=plp10025
3881.1	SEQID=6990	EMBL_NAME=plp10026
3882.1	SEQID=6991	EMBL_NAME=plp10027
3884.1	SEQID=6992	EMBL_NAME=plp10028
3886.2	SEQID=6993	EMBL_NAME=plpl0029
3888.1	SEQID=6994	EMBL_NAME=plpl0030
3890.1	SEQID=6995	EMBL_NAME=plpl0031
3891.2	SEQID=6996	EMBL_NAME=plp10032
3893.1	SEQID=6997	EMBL_NAME=plp10033
3894.1	SEQID=6998	EMBL_NAME=plpl0034
3949.1	SEQID=7002	EMBL NAME=lpl1158
3976.1	SEQID=7003	EMBL_NAME=lpl1138
3987.1	SEQID=7004	EMBL_NAME=lpl1132
4007.1	SEQID=7006	EMBL_NAME=lpl1116
4172.1	SEQID=7014	EMBL_NAME=lpl0194
4173.1	SEQID=7015	EMBL_NAME=lpl0193
4175.1	SEQID=7017	EMBL_NAME=lpl0191
4181.1	SEQID=7019	EMBL_NAME=lpl0189
4182.1	SEQID=7020	EMBL_NAME=lpl0188
4185.1	SEQID=7021	EMBL NAME=lpl0187
1189.2	SEQID=7023	EMBL NAME=lpl1424
1196.1	SEQID=7024	EMBL NAME=lpl1417
1197.1	SEQID=7025	EMBL_NAME=lpl1416
1248.1	SEQID=7030	EMBL_NAME=lpl1942
250.1	SEQID=7031	EMBL_NAME=lpl1943
251.1	SEQID=7032	EMBL_NAME=lpl1944
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4252.1	SEQID=7033	EMBL NAME=lpl1945
4253.1	SEQID=7034	EMBL NAME=lpl1946
4254.1	SEQID=7035	EMBL NAME=lpl1947
560.1	SEQID=7037	EMBL NAME=lpl1681
561.1	SEQID=7037 SEQID=7038	
	•	EMBL_NAME=lpl1680
562.1	SEQID=7039	EMBL_NAME=lpl1679
564.1	SEQID=7040	EMBL_NAME=lpl1678
688.1	SEQID=7042	EMBL NAME=lpl1588
689.1	SEQID=7043	EMBL_NAME=lpl1587
692.1	SEQID=7045	EMBL NAME=lpl1585
699.1	SEQID=7048	EMBL_NAME=lpl1581
700.1	SEQID=7049	EMBL_NAME=lpl1580
703.1	SEQID=7050	EMBL_NAME=lpl1579
709.1	SEQID=7053	EMBL NAME=lpl1575
711.1	SEQID=7054	EMBL_NAME=lpl1574
760.1	SEQID=7055	EMBL NAME=lpl1537
761.1	SEQID=7056	EMBL NAME=lpl1536
898.1	SEQID=7057	EMBL_NAME=lpl1425
995.2	SEQID=7059	EMBL_NAME=lpl1933
997.1	SEQID=7060	EMBL_NAME=lpl1931
2144.1	SEQID=6820	EMBL_NAME=lpl1043b

<u>Table XIX</u>: List of the sequences present in the Paris and Lens strain though absent in the Philadelphia strain with their Pasteur Institute « ORF » correspondence number and accession number in the gene banks

1082.3	SEQID=3560	EMBL_NAME=lpp1844
1090.1	SEQID=3565	EMBL_NAME=lpp1061
1156.3	SEQID=3606	EMBL_NAME=lpp1106
119.1	SEQID=3625	EMBL NAME=lpp2953
121.1	SEQID=3635	EMBL NAME=lpp2952
1225.2	SEQID=6596	EMBL_NAME=plpp0013
1226.2	SEQID=6597	EMBL NAME=plpp0012
131.2	SEQID=3694	EMBL_NAME=lpp1099
1469.4	SEQID=3786	EMBL_NAME=lpp1843
15.1	SEQID=3803	EMBL_NAME=lpp0194
1560.2	SEQID=3842	EMBL_NAME=lpp2477
16.1	SEQID=3869	EMBL_NAME=lpp0193
1737.2	SEQID=3956	EMBL_NAME=lpp1863
1875.2	SEQID=4045	EMBL_NAME=lpp2529
2.1	SEQID=4112	EMBL_NAME=lpp0163
2026.1	SEQID=4124	EMBL_NAME=lpp1909
2039.1	SEQID=4131	EMBL_NAME=lpp0243
2275.4	SEQID=6632	EMBL_NAME=plpp0014
2357.4	SEQID=4299	EMBL_NAME=lpp2054
2427.4	SEQID=4348	EMBL_NAME=lpp1869
2453.4	SEQID=4367	EMBL_NAME=lpp2450
2649.1	SEQID=4483	EMBL_NAME=lpp0667
321.3	SEQID=4804	EMBL_NAME=lpp2981
3248.1	SEQID=4829	EMBL_NAME=lpp2478
33.1	SEQID=4861	EMBL_NAME=lpp0183
3395.3	SEQID=4911	EMBL_NAME=lpp1042
3396.1	SEQID=4912	EMBL_NAME=lpp1043
3401.2	SEQID=4917	EMBL_NAME=lpp1047
341.6	SEQID=4921	EMBL_NAME=lpp0024
3413.2	SEQID=4923	EMBL NAME=1pp2060
3414.3	SEQID=4924	EMBL_NAME=lpp2061
3499.1	SEQID=6673	EMBL_NAME=plpp0011
3500.3	SEQID=6674	EMBL_NAME=plpp0010
3563.3	SEQID=5023	EMBL_NAME=lpp0158
3594.1	SEQID=5041	EMBL_NAME=lpp0639
3600.2	SEQID=5048	EMBL_NAME=lpp2449
3601.1	SEQID=5049	EMBL_NAME=lpp2448
3657.1	SEQID=5085	EMBL_NAME=lpp2419
3734.1	SEQID=5134	EMBL_NAME=lpp0208
3744.2	SEQID=5142	EMBL_NAME=lpp1850
3763.1	SEQID=5152	EMBL_NAME=lpp1867
3871.1	SEQID=5211	EMBL_NAME=lpp2070
3872.1	SEQID=5212	EMBL_NAME=lpp2069
3878.1	SEQID=5215	EMBL_NAME=lpp2066

4039.2	SEQID=5315	EMBL NAME=lpp0064
4040.1	SEQID=5316	EMBL NAME=lpp0063
4045.2	SEQID=5320	EMBL NAME=lpp0059
417.3	SEQID=5412	EMBL NAME=lpp1907
4276.2	SEQID=5479	EMBL_NAME=lpp2048
4532.2	SEQID=5653	EMBL NAME=lpp2049
4763.2	SEQID=5803	EMBL NAME=lpp1088
4764.2	SEQID=5804	EMBL_NAME=lpp1087
5056.3	SEQID=5980	EMBL NAME=lpp1578
5058.2	SEQID=5981	EMBL_NAME=lpp1579
5059.3	SEQID=5982	EMBL_NAME=lpp1580
506.3	SEQID=5983	EMBL_NAME=lpp2417
5080.6	SEQID=6686	EMBL_NAME=plpp0006
5087.2	SEQID=5999	EMBL_NAME=lpp2016
5106.3	SEQID=6688	EMBL_NAME=plpp0007
5147.1	SEQID=6691	EMBL_NAME=plpp0009
5176.1	SEQID=6026	EMBL_NAME=lpp0712
5382.1	SEQID=6107	EMBL_NAME=lpp1086
5388.1	SEQID=6110	EMBL_NAME=lpp0084
5404.2	SEQID=6117	EMBL_NAME=lpp2443
5504.4	SEQID=6163	EMBL_NAME=lpp2053
553.1	SEQID=6173	EMBL_NAME=lpp2920
5584.2	SEQID=6200	EMBL_NAME=lpp1450
5609.1	SEQID=6214	EMBL_NAME=lpp2153
58.1	SEQID=6273	EMBL_NAME=lpp0165
6036.1	SEQID=6322	EMBL_NAME=lpp1449
650.4	SEQID=6385	EMBL_NAME=lpp0668
651.3	SEQID=6386	EMBL_NAME=lpp0669
860.2	SEQID=6495	EMBL_NAME=lpp2058
9.2	SEQID=6521	EMBL_NAME=lpp0159

<u>Table XX</u>: List of the sequences present in the Paris and Philadelphia strain though absent in the Lens strain with their Pasteur Institute « ORF » correspondence number and accession number in the gene banks

102.1	SEQID=3519	EMBL NAME=lpp2364
103.1	SEQID=3525	EMBL_NAME=lpp2363
104.1	SEQID=3533	EMBL_NAME=lpp2362
107.1	SEQID=3553	EMBL_NAME=lpp2361
109.1	SEQID=3564	EMBL_NAME=lpp2360
1107.2	SEQID=3578	EMBL_NAME=lpp1601
1109.3	SEQID=3579	EMBL_NAME=lpp1600
111.2	SEQID=3580	EMBL_NAME=lpp2359
1111.3	SEQID=3581	EMBL_NAME=lpp1599
1211.3	SEQID=3637	EMBL_NAME=lpp0258
1236.2	SEQID=6599	EMBL NAME=plpp0120
1334.3	SEQID=3707	EMBL_NAME=lpp0094
1335.2	SEQID=3708	EMBL_NAME=lpp0095
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1423.2	SEQID=3763	EMBL_NAME=lpp2328
1424.2	SEQID=3764	EMBL_NAME=lpp2329
1425.2	SEQID=3765	EMBL_NAME=lpp2330
1432.3	SEQID=3767	EMBL_NAME=lpp2441
152.3	SEQID=3818	EMBL NAME=Inp2354
1548.2	SEQID=3834	EMBL_NAME=lpp2331
1549.1	SEQID=3835	EMBL_NAME=lpp2332
1550.2	SEQID=3836	EMBL_NAME=lpp2333
158.2	SEQID=3854	EMBL_NAME=lpp2341
159.1	SEQID=3862	EMBL_NAME=lpp2342
160.1	SEQID=3870	EMBL_NAME=lpp2343
1628.3	SEQID=3885	EMBL_NAME=lpp2339
1629.1	SEQID=3886	EMBL_NAME=lpp2338
1631.3	SEQID=3888	EMBL_NAME=lpp2337
1635.4	SEQID=3891	EMBL_NAME=lpp1309
1639.4	SEQID=3894	EMBL_NAME=lpp1308
168.1	SEQID=3920	EMBL_NAME=lpp2346
1682.4	SEQID=3922	EMBL_NAME=lpp0045
169.1	SEQID=3927	EMBL_NAME=lpp2347
1703.1	SEQID=3936	EMBL_NAME=lpp1942
1775.3	SEQID=3983	EMBL_NAME=lpp1130
1847.3	SEQID=4029	EMBL_NAME=lpp1089
1887.2	SEQID=4053	EMBL_NAME=lpp1940
190.3	SEQID=4062	EMBL_NAME=lpp0234
1960.2	SEQID=4096	EMBL_NAME=lpp2340
1961.3	SEQID=4097	EMBL_NAME=lpp2355
2031.2	SEQID=4128	EMBL_NAME=lpp0330
2054.2	SEQID=4138	EMBL_NAME=lpp2603
2079.2	SEQID=4153	EMBL_NAME=lpp2455
2143.5	SEQID=4186	EMBL_NAME=lpp2615
2169.2	SEQID=4205	EMBL_NAME=lpp1890
227.2	SEQID=4247	EMBL_NAME=lpp2358
228.1	SEQID=4253	EMBL_NAME=lpp2357
2544.3	SEQID=4425	EMBL_NAME=lpp2192
2591.3	SEQID=4451	EMBL_NAME=lpp2779
2637.1	SEQID=4474	EMBL_NAME=lpp2336
2639.1	SEQID=4475	EMBL_NAME=lpp2327
2646.1	SEQID=4480	EMBL_NAME=lpp0673
2730.1	SEQID=4526	EMBL_NAME=lpp0251
2808.1	SEQID=4572	EMBL_NAME=lpp0321
2849.1	SEQID=4596	EMBL_NAME=lpp1912
2938.4	SEQID=4636	EMBL_NAME=lpp2883
2991.1	SEQID=4675	EMBL_NAME=lpp2110
2992.2	SEQID=4676	EMBL_NAME=lpp2109
3163.1	SEQID=4778	EMBL_NAME=lpp0331
3190.1	SEQID=4795	EMBL_NAME=lpp1007
3191.1	SEQID=4796	EMBL_NAME=lpp1006
3205.3	SEQID=4802	EMBL_NAME=lpp1006 EMBL_NAME=lpp2131
3207.5	SEQID=4803	EMBL_NAME=lpp2132
3250.1	SEQID=4832	EMBL_NAME=lpp2474
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3508.2	SEQID=4982	EMBL_NAME=lpp2780
3588.2	SEQID=5036	EMBL_NAME=lpp1090
3663.1	SEQID=5090	EMBL_NAME=lpp3017
3699.3	SEQID=5114	EMBL_NAME=lpp1404
3701.1	SEQID=5116	EMBL_NAME=lpp1403
371.1	SEQID=5123	EMBL_NAME=lpp2039
3783.1	SEQID=5160	EMBL_NAME=lpp1144
3822.2	SEQID=5187	EMBL_NAME=lpp0851
384.5	SEQID=5197	EMBL_NAME=lpp0716
3884.2	SEQID=5216	EMBL NAME=lpp2440
402.2	SEQID=5304	EMBL NAME=lpp1557
4030.1	SEQID=5310	EMBL_NAME=lpp0096
4083.4	SEQID=5348	EMBL_NAME=lpp1603
4084.3	SEQID=5349	EMBL_NAME=lpp1602
4220.2	SEQID=5443	EMBL_NAME=lpp1852
423.2	SEQID=5449	EMBL_NAME=lpp1832
424.1	SEQID=5457	EMBL_NAME=lpp2431
4242.3	•	EMBL_NAME=1pp2431
4249.1	SEQID=5458	EMBL_NAME=lpp1405
4453.2	SEQID=5464	EMBL_NAME=lpp3009
4517.2	SEQID=5597	EMBL_NAME=lpp1447
	SEQID=5640	EMBL_NAME=lpp2497
4528.2	SEQID=5649	EMBL_NAME=lpp2052
4530.1	SEQID=5651	EMBL_NAME=lpp2051
4559.1	SEQID=5667	EMBL_NAME=lpp0287
4591.2	SEQID=5689	EMBL_NAME=lpp2508
4819.2	SEQID=5845	EMBL_NAME=lpp2498
4965.2	SEQID=5926	EMBL_NAME=lpp0829c
4966.2	SEQID=5927	EMBL_NAME=lpp0830
5060.2	SEQID=5984	EMBL_NAME=lpp0835
5289.2	SEQID=6070	EMBL_NAME=lpp0589
5328.1	SEQID=6088	EMBL_NAME=lpp1565
5337.1	SEQID=6090	EMBL_NAME=lpp2896
5340.1	SEQID=6091	EMBL_NAME=lpp1947
538.1	SEQID=6105	EMBL_NAME=lpp0859
5496.2	SEQID=6157	EMBL_NAME=lpp0829b
5656.2	SEQID=6238	EMBL_NAME=lpp2887
5723.2	SEQID=6261	EMBL_NAME=lpp1944
5871.4	SEQID=6288	EMBL_NAME=lpp2037
590.4	SEQID=6299	EMBL_NAME=lpp2886
592.3	SEQID=6305	EMBL_NAME=lpp2348
5920.3	SEQID=6306	EMBL_NAME=lpp2311
593.2	SEQID=6308	EMBL_NAME=lpp2349
594.1	SEQID=6309	EMBL_NAME=lpp2350
5999.1	SEQID=6317	EMBL_NAME=lpp0039
6002.2	SEQID=6318	EMBL_NAME=lpp0881
6110.1	SEQID=6334	EMBL_NAME=lpp0829a
615.5	SEQID=6337	EMBL_NAME=lpp1562
6151.1	SEQID=6338	EMBL NAME=lpp0038
6159.1	SEQID=6339	EMBL_NAME=lpp2410
6160.1	SEQID=6341	EMBL NAME=lpp2402

6178.1	SEQID=6343	EMBL_NAME=lpp0210
6180.1	SEQID=6345	EMBL_NĂME=lpp1035
6186.1	SEQID=6346	EMBL_NAME=lpp0882
6195.2	SEQID=6350	EMBL_NAME=lpp0717
6285.1	SEQID=6362	EMBL_NAME=lpp2895
6309.1	SEQID=6364	EMBL_NAME=lpp1945
6318.1	SEQID=6366	EMBL_NAME=lpp1851
6320.1	SEQID=6368	EMBL_NAME=lpp1813
6322.1	SEQID=6369	EMBL_NAME=lpp1812
743.4	SEQID=6432	EMBL_NAME=lpp2368
744.4	SEQID=6433	EMBL_NAME=lpp2369
818.2	SEQID=6469	EMBL_NAME=lpp2335
819.2	SEQID=6470	EMBL_NAME=lpp2334
864.1	SEQID=6498	EMBL_NAME=lpp2055
901.2	SEQID=6525	EMBL_NAME=lpp2471
938.3	SEQID=6550	EMBL_NAME=lpp1253
96.6	SEQID=6562	EMBL_NAME=lpp2367
97.2	SEQID=6568	EMBL_NAME=lpp2366
979.3	SEQID=6574	EMBL_NAME=lpp2494
980.1	SEQID=6575	EMBL_NAME=lpp2495
981.2	SEQID=6576	EMBL_NAME=lpp2496
	•	

<u>Table XXI</u>: List of the sequences present in the Philadelphia and Lens strain though absent in the Paris strain with their Pasteur Institute « ORF » correspondence number and accession number in the gene banks

1038.1 SEOID=6741 EMBL NAME=lpl1901 1043.1 SEQID=6744 EMBL NAME=lpl1897 1073.1 SEQID=6753 EMBL NAME=lpl0044 1130.1 SEQID=6759 EMBL NAME=lpl2933 117.1 SEQID=6762 EMBL NAME=lpl0171 124.1 SEQID=6771 EMBL NAME=lp10167 125.1 SEQID=6772 EMBL NAME=lpl0166 1282.1 SEQID=6785 EMBL NAME=lpl2836 1434.1 SEOID=6793 EMBL NAME=lpl2732 148.1 SEQID=6794 EMBL NAME=lp10150 175.1 SEQID=6802 EMBL NAME=lpl0132 1825.1 SEQID=6804 EMBL NAME=lp10802 1828.1 SEQID=6807 EMBL NAME=lp10805 1829.1 SEQID=6808 EMBL NAME=lpl0806 2100.1 SEQID=6811 EMBL NAME=lpl1006 2134.1 SEQID=6813 EMBL NAME=lpl1034 2135.1 SEQID=6814 EMBL NAME=lpl1035 2136.1 SEQID=6815 EMBL NAME=lpl1036 2139.1 SEQID=6817 EMBL NAME=lpl1039 2151.1 SEQID=6822 EMBL NAME=lpl1047 2157.1 SEQID=6825 EMBL NAME=lpl1051 2168.1 SEQID=6826 EMBL NAME=lpl1058

2102.1	CECTO 1	
2192.1	SEQID=6834	EMBL_NAME=lpl1075
2205.1	SEQID=6843	EMBL NAME=lpl1086
2209.1	SEQID=6847	EMBL_NAME=lpl1090
2238.1	SEQID=6852	EMBL_NAME=lpl1110
2244.1	SEQID=6855	EMBL_NAME=lpl0201
2246.1	SEQID=6857	EMBL_NAME=lpl0203
2247.1	SEQID=6858	EMBL_NAME=lpl0204
2248.1	SEQID=6859	EMBL_NAME=lpl0205
2261.1	SEQID=6863	EMBL_NAME=lpl0213
2392.1	SEQID=6864	EMBL_NAME=lpl2580
2422.1	SEQID=6865	EMBL_NAME=lpl2558
2517.1	SEQID=6869	EMBL_NAME=lpl2496
2539.1	SEQID=6883	EMBL_NAME=lpl2478
2558.1	SEQID=6887	EMBL_NAME=lpl2465
2587.1	SEQID=6889	EMBL_NAME=lpl2443
2660.1	SEQID=6893	EMBL_NAME=lp12384
2696.1	SEQID=6896	EMBL_NAME=lpl2358
2706.1	SEQID=6899	
2777.1	SEQID=6911	EMBL_NAME=1p12351
2807.1	SEQID=6920	EMBL_NAME=1p12308
2809.1	SEQID=6922	EMBL_NAME=1p12287
293.1	SEQID=6924	EMBL_NAME=lpl2285
3151.1	SEQID=6934	EMBL_NAME=1,120.42
322.1	SEQID=6941	EMBL_NAME=lpl2042
3536.1	SEQID=6946	EMBL_NAME=lpl1857
3537.1	SEQID=6947	EMBL_NAME=lpl0286
3749.2	SEQID=6954	EMBL_NAME=lp10287
3788.1	SEQID=6955	EMBL_NAME=lpl0569
3793.1	SEQID=6956	EMBL_NAME=lpl0593
3816.1		EMBL_NAME=lpl0596
3827.1	SEQID=6958	EMBL_NAME=plp10038
3828.1	SEQID=6963	EMBL_NAME=plpl0044
3831.1	SEQID=6964	EMBL_NAME=plpl0045
3835.1	SEQID=6965	EMBL_NAME=plpl0046
3836.1	SEQID=6966	EMBL_NAME=plpl0047
3837.1	SEQID=6967	EMBL_NAME=plpl0048
3839.1	SEQID=6968	EMBL_NAME=plpl0049
3840.1	SEQID=6969	EMBL NAME=plp10050
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	SEQID=6975	EMBL_NAME=plpl0011
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3862.1	SEQID=6977	EMBL_NAME=plpl0013
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3937.1	SEQID=7001	EMBL_NAME=lpl1165
3995.1	SEQID=7005	EMBL_NAME=lpl1125
4011.1	SEQID=7007	EMBL NAME=lpl1113
4085.1	SEQID=7008	EMBL NAME=1p10408
4093.1	SEQID=7009	EMBL_NAME=lpl0415
4111.1	SEQID=7010	EMBL_NAME=lpl0432

4169.1	SEQID=7011	EMBL_NAME=lpl0197
4170.1	SEQID=7012	EMBL_NAME=lpl0196
4171.2	SEQID=7013	EMBL_NAME=lpl0195
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4188.1	SEQID=7022	EMBL_NAME=lp10185
4203.1	SEQID=7026	EMBL_NAME=lpl1412
4229.2	SEQID=7027	EMBL_NAME=lpl1393
4236.2	SEQID=7028	EMBL_NAME=lpl1965
4237.1	SEQID=7029	EMBL_NAME=lpl1934
442.1	SEQID=7036	EMBL_NAME=lpl1768
566.1	SEQID=7041	EMBL_NAME=lpl1676
690.1	SEQID=7044	EMBL_NAME=lpl1586
694.1	SEQID=7046	EMBL_NAME=lpl1584
697.1	SEQID=7047	EMBL_NAME=lpl1582
705.1	SEQID=7051	EMBL_NAME=lpl1578
707.1	SEQID=7052	EMBL_NAME=lpl1577
899.2	SEQID=7058	EMBL NAME=In12032

Legionella specific	•	ı	ı			•	+	+			•	•	+	1	•	+	+	ı	,	· 1	1	ı	ı	+	1	•	•	ı	Ī	•
Presence in L. longbeachae																								+						
% homology retained ORFs		%26																94%	%86	%26	%66	100%	%66	%66	%86	%96	%26	100%	100%	100%
Specific to Paris/ Philadelphia	+	•	+	+	+	+	+	+	+	+	+	+	+	+	+	+	+	,	1	•	,	•	•	1	ı	•	•		1	
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	541	1703	2313	1048	2223	3074	3130	3457	220	1391	2307	2898	3113	4808	5936	6501	862	2051	3128	3735	4526	4866	5745	6974	7888	8814	10349	290	1026	2307
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ORF	5906.1	5905.1	6131.1	6130.1	6128.1	6125.1	6123.1	6122.1	6121.1	3257.2	3258.2	1924.4	1923.2	1920.3	5626.2	5987.1	5898.2	523.2	524.2	526.2	528.2	529.2	3609.1	3610.1	3611.2	3612.2	3613.3	3496.2	3494.2	216.2
SEQ ID No.	SEQ ID No. 3098	SEQ ID No. 3097	SEQ ID No. 3181	SEQ ID No. 3180	SEQ ID No. 3179	SEQ ID No. 3178	ò	SEQ ID No. 3176	SEQ ID No. 3175	SEQ ID No. 1494	SEQ ID No. 1495	SEQ ID No. 659	SEQ ID No. 658	SEQ ID No. 657	SEQ ID No. 2987	SEQ ID No. 3122	D No.	SEQ ID No. 2792	ġ	SEQ ID No. 2804	SEQ ID No. 2812	SEQ ID No. 2816	SEQ ID No. 1736	SEQ ID No. 1737	SEQ ID No. 1738	SEQ ID No. 1739	SEQ ID No. 1740	SEQ ID No. 1651	ġ	SEQ ID No. 789

100% 100% 100% 100% 100% 100% 100% 99% 100% 100% %66 %66 %66 91% 79% 94% 100% 89% 87% 92% 87% 20884 21334 21738 22130 22130 222697 23012 23012 23012 234357 24873 25495 2549 20459
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6445
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8412 Contig37
Contig37 Contig39 Contig39 Contig39 Contig39 Contig39 4133.1 4133.1 4134.1 4136.1 4136.1 4140.1 4144.3 41 SEQ ID No. 2073
SEQ ID No. 2075
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SEQ ID No. 2077
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SEQ ID No. 2080
SEQ ID No. 2081
SEQ ID No. 2081
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SEQ ID No. 2080
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SEQ ID No. 3087
SEQ ID No. 3088
SEQ ID No. 1965
SEQ ID No. 1586
SEQ ID No. 1589

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10196	10565	11457	12883	13558	13921	14302	14931	16069	18045	18568	20124	21501	22468	25228	25532	26788	27551	28200	28574	29202	29538	30161	30592	31406	31603	31995	32401	32613	32977	34046	34304	35343	36188	36741	37811
٥	Ε	Ε	۵	Ε	ε	۵	Ε	٤	٤	Ε	Ε	Ε	Ε	Ε	٤	E	E	٤	Ε	٤	٤	Ε	٤	٤	E	E	ε	٤	Ε	a	۵	Ε	٤	٤	E
Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39	Contig39									
3401.2	3402.2	3404.2	3406.2	5322.1	5321.1	4914.2	4913.2	9.779	675.6	673.3	3575.2	3574.1	1089.2	1090.1	1091.2	1092.3	5031.2	5032.1	5033.1	5319.1	5318.2	4132.3	4131.3	5776.1	4130.2	4129.1	4128.1	4127.1	4126.2	4123.2	682.2	681.3	680.3	678.3	4955.4
SEQ ID No. 1593	SEQ ID No. 1594	SEQ ID No. 1595	SEQ ID No. 1596	SEQ ID No. 2832	SEQ ID No. 2831	SEQ ID No. 2615	SEQ ID No. 2614	SEQ ID No. 3226	SEQ ID No. 3225	№	В В		SEQ ID No. 116	ID No.	SEQ ID No. 119	SEQ ID No. 120	SEQ ID No. 2684	SEQ ID No. 2685	SEQ ID No. 2686	SEQ ID No. 2829	SEQ ID No. 2828	SEQ ID No. 2072	SEQ ID No. 2071	Ω So		№	Ю Мо		Ю В	Ö. Se			D No	Ю Мо	SEQ ID No. 2639

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39111 39610	40214	40834	41153	41667	43268	45192	46123	47244	48300	49143	49497	20090	50780	51193	51668	52705	56130	57019	57909	58368	59289	60441	61850	2	534	2006	3481	5934	6795	8321	10617	11840	13158	15227
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4954.2 4952.1	4951.2	5382.1	4764.2	4763.2	1847.3	3588.2	3586.2	1229.3	1227.1	2383.4	1390.5	1391.1	1392.2	3584.1	131.2	6117.1	577.2	3729.1	3730.1	3731.1	3732.1	3734.1	3735.2	5874.1	615.5	2767.1	399.2	400.1	401.2	402.2	2120.2	1075.3	1074.3	2121.2
SEQ ID No. 2638 SEQ ID No. 2637	ID No.	SEQ ID No. 2858	SEQ ID No. 2515	SEQ ID No. 2514	SEQ ID No. 609	SEQ ID No. 1718	SEQ ID No. 1717	SEQ ID No. 207	SEQ ID No. 206	D No.	SEQ ID No. 309	SEQ ID No. 310	SEQ ID No. 311	SEQ ID No. 1716	SEQ ID No. 260	SEQ ID No. 3172	SEQ ID No. 3047	SEQ ID No. 1816	SEQ ID No. 1817	SEQ ID No. 1818	SEQ ID No. 1819	SEQ ID No. 1820	SEQ ID No. 1821	SEQ ID No. 3079	SEQ ID No. 3185	SEQ ID No. 1176	SEQ ID No. 1969	SEQ ID No. 1978	SEQ ID No. 1985	SEQ ID No. 1993	SEQ ID No. 760	SEQ ID No. 107	SEQ ID No. 106	SEQ ID No. 761

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SEQ ID No. 1175	2765.2	Contig40	Ε	16013	16903		ı	95%	+
SEQ ID No. 1174	2763.2	Contig40	Ε	16914	17111		+		+
SEQ ID No. 1173	2761.1	Contig40	۵	17411	17758	1	1	100%	
SEQ ID No. 1172	2760.1	Contig40	. Δ	17730	17999	ı	+		•
SEQ ID No. 1170	2759.1	Contig40	a	17938	18912	1	ı	%86	•
SEQ ID No. 1169	2757.1	Contig40	۵	18913	19374	1	•	%86	
SEQ ID No. 1168	2755.1	Contig40	۵	19564	21600	ı	•	. %86	+
SEQ ID No. 1167	2754.2	Contig40	٤	21696	22277	1	ı	%66	,
SEQ ID No. 1166	2753.1	Contig40	Ε	22278	22781		•	%86	+
SEQ ID No. 1165	2752.1	Contig40	٤	22782	23303	+	•	100%	,
SEQ ID No. 1164	2751.1	Contig40	Ε	23593	24363		•	%66	ı
SEQ ID No. 3351	859.2	Contig40	Q	24703	26091		•	+ %66	,
SEQ ID No. 3350	858.2	Contig40	۵	26045	26566		•	%26	•
SEQ ID No. 371	1493.6	Contig40	۵	26618	31495		ı	+ %66	•
SEQ ID No. 509	1697.2	Contig40	ď	31501	32064	1		%86	
SEQ ID No. 510	1698.2	Contig40	Ε	32095	33180		•	%96	ı
SEQ ID No. 2882	5429.1	Contig40	ο.	33195	33449			%46	•
SEQ ID No. 1047	2562.2	Contig40	a	33385	33834			95%	+
SEQ ID No. 1046	2560.3	Contig40	ď	33902	34498			%66	•
SEQ ID No. 1044	2559.3	Contig40	۵	34609	35655		•	%66	•
SEQ ID No. 2881	5428.1	Contig40	a	35656	36054			100%	•
SEQ ID No. 2880	5427.1	Contig40	۵	36055	36756	,	ı	100%	•
SEQ ID No. 2879	5426.2	Contig40	۵	36750	37220	1	,	100%	
SEQ ID No. 1010	2504.4	Contig40	ď	37208	38215	,	1	%66	
SEQ ID No. 1009	2501.2	Contig40	۵	38209	39480	+	•	%66	•
SEQ ID No. 1572	3367.1	Contig40	۵	39438	40316		•	%86	•
SEQ ID No. 3241	699.3	Contig40	Ε	40345	41484	1	ı	%96	
SEQ ID No. 3243	701.2	Contig40	Ε	41465	42997	1	ı	%96	•
SEQ ID No. 3244	702.2	Contig40	Ε	42928	43755	1	ı	%96	
SEQ ID No. 1571	3365.2	Contig40	E	43915	44994		ı	%96	•
SEQ ID No. 1570	3363.2	Contig40	E	44988	45944		ı	%66	
SEQ ID No. 1569	3362.1	Contig40	a	46177	46929			%26	
ġ	3360.1	Contig40	۵	46986	47717	+		%68	+
<u>_</u>	1190.4	Contig40	E	47773	49917	+	ı	%96	
Θ S	3359.2	Contig40	Ε	20090	52309		ı .	%66	
SEQ ID No. 3026	5701.2	Contig40	٤	52481	53623	1	ı	%66	ı

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56410 Contig41 341.6 5380.3 4395.2 4396.1 4398.2 5644.3 4919.1 4918.2 5642.1 3895.2 3895.2 3895.2 3895.3 1041.3 1041.3 2404.2 2202.3 1174.2 1174.2 2815.2 2816.2 2816.2 SEQ ID No. 2856
SEQ ID No. 2263
SEQ ID No. 2264
SEQ ID No. 2265
SEQ ID No. 2996
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SEQ ID No. 3373
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SEQ ID No. 938
SEQ ID No. 2689
SEQ ID No. 173
SEQ ID No. 1205
SEQ ID No. 1207
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62316	62699	63314	63867	64548	64932	65284	60299	66491	66699	67559	69211	70081	71460	72529	74894	75964	78109	79904	82606	297	831	1367	2210	2533	4196	5536	0029	8247	9150	10583	11344	12007	12639	13730	14850
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2127.2	2129.1	2130.1	2132.2	2133.1	2135.1	2136.1	2137.1	2138.2	2140.2	1552.3	1555.2	1556.2	2824.1	910.4	909.2	2119.2	2826.1	2827.3	2828.3	5521.1	5520.1	3684.2	3685.1	3686.1	3687.1	3688.1	776.3	778.2	779.4	4119.2	4118.3	4116.3	4115.1	1764.3	1765.4
SEQ ID No. 763	SEQ ID No. 764	SEQ ID No. 766	SEQ ID No. 767	SEQ ID No. 768	SEQ ID No. 769	SEQ ID No. 770	SEQ ID No. 771	SEQ ID No. 772	SEQ ID No. 774	SEQ ID No. 412	SEQ ID No. 413	SEQ ID No. 414	SEQ ID No. 1211	SEQ ID No. 3389	SEQ ID No. 3388	SEQ ID No. 758	SEQ ID No. 1212	SEQ ID No. 1213	SEQ ID No. 1214	SEQ ID No. 2927	SEQ ID No. 2926	SEQ ID No. 1793	SEQ ID No. 1794	SEQ ID No. 1795	SEQ ID No. 1796	SEQ ID No. 1797	SEQ ID No. 3298	SEQ ID No. 3299	SEQ ID No. 3300	SEQ ID No. 2062	SEQ ID No. 2061	SEQ ID No. 2060	ġ	Š	SEQ ID No. 556

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					ġ.	SEQ ID No. 2916	ġ	ġ	SEQ ID No. 2646	ġ	ġ	ġ		SEQ ID No. 2932	SEQ ID No. 2705	SEQ ID No. 2706	SEQ ID No. 2091	SEQ ID No. 2092	SEQ ID No. 2093	D No.	SEQ ID No. 287	Ю В	D No	D No	ID No.	ID No	Θ S S	SEQ ID No. 977	Ю No	SEQ ID No. 1875	SEQ ID No. 1874	₽	ID No.	SEQ ID No. 879

SEQ ID No. 974	2450.3	Contig43	E		54343	1	1	%86	
SEQ ID No. 340	1438.4	Contig43	۵		56333		ı	%86	
ID No.	521.2	Contig43	E		58832		•	%86	ı
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SEQ ID No. 1234	2868.1	Contig43	۵		63514	+	•	%86	
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SEQ ID No. 1052	2570.2	Contig43	Ε		66153	+	1	%66	
SEQ ID No. 1053	2571.2	Contig43	٤		67073		1	%26	
SEQ ID No. 1232	2863.1	Contig43	E		67788	,		94%	
SEQ ID No. 1231	2862.1	Contig43	٤		68956	1	1	%96	
SEQ ID No. 156	1148.2	Contig43	E		71929			%86	,
SEQ ID No. 541	1746.2	Contig43	a		72792		1	91%	
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SEQ ID No. 1230	2861.1	Contig43	۵		74438	•	1	%26	•
SEQ ID No. 969	2443.2	Contig43	Д		75246		1	95%	+
SEQ ID No. 968	2442.2	Contig43	Ε		76591	1		%66	
SEQ ID No. 2502	4746.1	Contig43	٤		76862	+		91%	
SEQ ID No. 2503	4748.1	Contig43	٤		77648		•	%66	
SEQ ID No. 2504	4749.2	Contig43	٤		9966		1	%86	ı
SEQ ID No. 1442	3172.2	Contig43	٤		80918			%66	ı
SEQ ID No. 3066	5824.1	Contig43	E		81201		+		
SEQ ID No. 1443	3173.1	Contig43	۵.		81654	+	ı	%86	•
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SEQ ID No. 1446	3177.1	Contig43	۵.		83762		ı	%86	ı
SEQ ID No. 1447	3178.1	Contig43	۵		84629			%86	1
SEQ ID No. 1448	3179.1	Contig43	<u>م</u>		86452	+	•	%66	
D No.	6107.1	Contig43	٤		86489		+		+
ID No.	1314.3	Contig43	E		88240	+		%66	ı
SEQ ID No. 1450	3181.2	Contig43	E		89379			%66	ı
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SEQ ID No. 3168	6106.1	Contig43	Ε		89972		+		
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1675.3	1673.5	1669.3	1670.3	4222.3	4221.3	4220.2	384.3	1685.3	914.3	915.3	5176.1	5177.2	5735.2	5598.2	6098.1	4387.3	4388.1	1044.3	1042.5	5600.2	588.3	587.2	585.2	583.2	6097.1	6094.1	4620.1	4618.1	4616.2	5672.1	4615.3	5479.2	3747.2	3748 1
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174.1	176.1	178.1	179.1	180.1	182.3	3008.1	2580.2	2579.2	3009.2	4546.3	2369.3	2371.1	2372.3	4542.4	1689.3	1691.3	1752.2	1750.3	2884.1	114.2	115.2	116.1	6083.1	117.1	118.1	119.1	121.1	5765.1	122.3	1979.1	1978.1	1976.2	2166.2	2167.2	2882.2
SEQ ID No. 537	SEQ ID No. 551	SEQ ID No. 566	SEQ ID No. 574	SEQ ID No. 581	SEQ ID No. 594		SEQ ID No. 1059	SEQ ID No. 1057	SEQ ID No. 1331	SEQ ID No. 2365	SEQ ID No. 916	SEQ ID No. 917	SEQ ID No. 918	SEQ ID No. 2364	SEQ ID No. 503	SEQ ID No. 505	SEQ ID No. 546	SEQ ID No. 545	SEQ ID No. 1243	SEQ ID No. 152	SEQ ID No. 157	SEQ ID No. 163	SEQ ID No. 3159		SEQ ID No. 176		SEQ ID No. 194	SEQ ID No. 3043	ID No.		Ö. No		ID No.		SEQ ID No. 1242

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126062	126066	126562	128457	128699	130078	131759	132141	133569	134694	136325	137595	137887	138934	139970	141531	142854	143330	143912	144919	145238	146551	148217	150016	150832	153153	153548	155278	157395	158250	161403	က	299	1499	1861	2147
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3053.3	6079.1	2070.4	1014.2	1013.1	2068.3	4669.2	4668.1	4666.2	1482.4	2034.2	5334.1	315.3	317.1	318.2	1812.4	2295.3	2297.3	4907.2	5337.1	5338.2	4266.3	723.3	722.3	719.4	5671.2	5669.2	5668.2	5656.2	590.4	4599.2	6074.1	4421.2	4422.1	4424.1	4425.1
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3129	3570	4176	2208	6970	7878	9075	9817	10206	10897	12027	12817	13726	14349	15500	16011	16733	17487	18508	19228	20119	21045	22114	22941	24389	25543	26936	28480	29682	30690	31594	32699	33556	21515
Ε (⊋ E	۵	۵.	Ε	٤	E	٤	٤	۵	Ε	a	Ε	٤	٤	E	E	E	E	a.	Ε	Ω.	۵	a	a	Q.	٤	۵	۵	٤	٥	٤	α.	c
Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contig49	Contina
4427.1	4429.2	1827.3	1826.2	4920.4	1302.2	1303.3	3570.2	2392.2	2391.2	3572.1	1578.3	1579.4	5044.2	5046.2	5047.3	2739.3	2738.2	2737.2	190.3	191.1	193.3	195.3	197.3	199.1	201.2	2030.2	2041.2	2039.1	494.2	496.1	497.1	498.4	5761 1
SEQ ID No. 2282		ġ	SEQ ID No. 596	SEQ ID No. 2621	SEQ ID No. 256	SEQ ID No. 257	SEQ ID No. 1709	SEQ ID No. 931	SEQ ID No. 930	SEQ ID No. 1710	SEQ ID No. 428	SEQ ID No. 429	SEQ ID No. 2694	SEQ ID No. 2695	SEQ ID No. 2696	SEQ ID No. 1156	SEQ ID No. 1155	SEQ ID No. 1154	SEQ ID No. 645	SEQ ID No. 651	SEQ ID No. 662	SEQ ID No. 675	SEQ ID No. 683	SEQ ID No. 691	SEQ ID No. 700	SEQ ID No. 712	SEQ ID No. 718	SEQ ID No. 716	SEQ ID No. 2629	SEQ ID No. 2642	SEQ ID No. 2650	SEQ ID No. 2655	SEO ID No. 3042

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SEQ ID No. 1153	2733.1	Contig49	Ω.	36823	37893	1		%68		,
SEQ ID No. 1152	2732.1	Contig49	٤	38090	38974	.+	ı	93%		+
SEQ ID No. 1151	2730.1	Contig49	۵	39326	40288	ı	•	%26		+
SEQ ID No. 182	1188.2	Contig49	Ε	40523	42844	+	1	%86		,
SEQ ID No. 663	1930.3	Contig49	E	42892	43923		•	94%		+
SEQ ID No. 720	2049.2	Contig49	٤	44183	45454	1	ı	%68		+
SEQ ID No. 1149	2728.1	Contig49	۵	45792	46145			%26		
S S	2727.2	Contig49	۵	46358	49045		+			.+
SEQ ID No. 197	1213.2	Contig49	Ф	49373	50596	+	+			
Ю В	1211.3	Contig49	a.	50962	52008			%98		
В В	930.3	Contig49	Ф	52100	53491		•	· %66		1
SEQ ID No. 3405	932.1	Contig49	۵	53437	54480	1		%86		
SEQ ID No. 3406	935.2	Contig49	٤	54481	55269		•	%66		
SEQ ID No. 3407	936.3	Contig49	Ε	55263	55808		ı	%26		+
SEQ ID No. 1177	2769.2	Contig49	E	55789	57639			%96		,
SEQ ID No. 1179	2770.1	Contig49	۵	57644	58357		ı	٠	_	,
SEQ ID No. 1180	2771.2	Contig49	ď	58350	59543		•	%26		ı
SEQ ID No. 3104	5926.1	Contig49	۵	59524	59913	,	+			,
SEQ ID No. 1181	2772.1	Contig49	Ε	59918	60340		•	%86		
SEQ ID No. 3214	657.3	Contig49	۵	60518	63580		,	%96		
SEQ ID No. 3213	656.3	Contig49	۵	63581	62269		,	94%		+
SEQ ID No. 1655	350.3	Contig49	Q.	65586	66236		•	%86		+
SEQ ID No. 1664	351.1	Contig49	٤	82299	66824		•	91%		
SEQ ID No. 1685	354.1	Contig49	Ε	66885	68309		ı	%26		
SEQ ID No. 1708	357.2	Contig49	E	68395	69129	ı	•	%86		
SEQ ID No. 1719	359.2	Contig49	E	08069	00/69		ı	82%		
SEQ ID No. 1182	2774.1	Contig49	α	69554	70459	1	ı	%26		
SEQ ID No. 1183	2775.1	Contig49	Q	70567	70794	1	+			
ġ	2777.1	Contig49	Ε	70880	71971	ı	1	94%		,
D No.	2778.2	Contig49	E	71956	72480	1	i	%26		,
Ю Мо	2780.2	Contig49	E	72516	73382		1	%26		
S S	2781.1	Contig49	Ε	73383	73757	1	ı	%66		
ID No.	1649.3	Contig49	Ε	73750	76110		1	94%		
D No.	1650.2	Contig49	E	76053	77585		ı	%96		
	2587.2	Contig49	σ	77654	78526	,	ı	%96		
SEQ ID No. 669	1940.2	Contig49	a	78565	80157	+	ı	%86		

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SEO ID No. 670	1943.3	Contig49	ď	80136	81140	+	•	+ %86	
SEQ ID No. 2375	4561.2	Contig49	ď	81131	82534		1	%86	•
Ö So	4560.2	Contig49	۵	82681	83823	,	ı	%26	+
SEQ ID No. 2373	4559.1	Contig49	٤	83994	85562	+	,	%86	•
SEQ ID No. 2372	4558.1	Contig49	٤	85707	86597	•	•	%26	+
SEQ ID No. 3155	6072.1	Contig49	۵	86701	86892		+		1
SEQ ID No. 3154	6071.1	Contig49	a	86822	87034		+		•
SEQ ID No. 2371	4554.2	Contig49	Ε	87282	87542		+		•
SEQ ID No. 2370	4553.2	Contig49	a	87651	88352	+	+		ı
SEQ ID No. 2752	5146.2	Contig49	q	88433	89740	+	+		•
SEQ ID No. 1402	3116.2	Contig49	٤	89715	90125		+		•
SEQ ID No. 1401	3114.2	Contig49	ď	90204	91163		+		- I,
D No.	3113.1	Contig49	ď	91157	93160		+		(1
SEQ ID No. 1399	3110.1	Contig49	۵	93148	93756	•	+		
SEQ ID No. 1398	3109.2	Contig49	a	93743	94096		+		
SEQ ID No. 1397	3107.2	Contig49	Ε	94191	94412		+		+
D No.	3106.2	Contig49	Ε	94616	96460		ı	%96	•
₽	3105.2	Contig49	Ε	96450	96755		+		
ID No.	3104.2	Contig49	٤	96892	99459		•	%26	
SEQ ID No. 1393	3102.2	Contig49	a	99671	100228			%86	•
Ω	3101.1	Contig49	۵	100453	102348		+		•
D No.	514.5	Contig49	Ε	102459	107018			. %26	+
D No.	436.4	Contig49	۵	104371	104940			%96	+
SEQ ID No. 783	2150.1	Contig49	٤	107367	107894		•	%26	•
Ю В	1222.5	Contig49	Ε	107944	109923			%96	
В В	2315.3	Contig49	۵	110157	110999		1	93%	۲.
SEQ ID No. 878	2317.2	Contig49	۵	111016	112482		ı	%26	
SEQ ID No. 1199	2804.1	Contig49	۵	112445	113809		1	%86	•
SEQ ID No. 858	2279.2	Contig49	۵	113883	114656		+		+
SEQ ID No. 525	1722.2	Contig49	۵	114908	115852		,	%26	. 1
SEQ ID No. 526	1723.5	Contig49	۵	115955	116539		1	%86	
SEQ ID No. 3153	6070.1	Contig49	а	116472	116879		+		
SEQ ID No. 1200	2806.1	Contig49	٤	117016	117618		,	%66	•
ID No.	670.1	Contig49	Ε	117945	119357			%26	•
SEQ ID No. 1385	309.2	Contig49	Ε	119391	122780		ı	%86	l'
SEQ ID No. 1391	310.1	Contig49	۵	122902	123675	•	ı	%96	• +

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5352	7022	8328	9435	12076	12476	13818	14561	15180	15646	15933	16229	16525	17281	17766	18538	19115	19555	19761	20260	21020	21804	22089	23502	24755	27710	27916	30127	30561	31477	31961	32483	33612	33985	34362	34254
5113	5355	6937	8329	9299	12084	12430	13740	14509	15176	15535	15924	16190	16526	17215	17675	18522	19109	19540	19757	20085	21133	21826	22255	23499	24756	27632	27917	30361	30803	31500	32298	32725	33710	34081	34102
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Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50
4233.3	1624.4	1623.4	4822.3	1287.3	5517.1	4104.2	4105.1	4106.1	4107.1	2367.2	1372.2	1373.1	1375.2	1376.2	4108.1	4109.2	5814.1	5813.1	5812.1	2625.2	2626.1	1803.2	1804.2	2627.1	2631.4	5953.1	747.3	5164.1	744.3	743.3	5952.1	9.96	97.2	99.1	5162.1
SEQ ID No. 2143	SEQ ID No. 459	SEQ ID No. 458	SEQ ID No. 2561	SEQ ID No. 247	SEQ ID No. 2925	SEQ ID No. 2050	SEQ ID No. 2051	SEQ ID No. 2052	SEQ ID No. 2053	SEQ ID No. 914	SEQ ID No. 297	SEQ ID No. 298	SEQ ID No. 299	SEQ ID No. 300	SEQ ID No. 2054	SEQ ID No. 2055	SEQ ID No. 3060	SEQ ID No. 3059	SEQ ID No. 3058	SEQ ID No. 1086	SEQ ID No. 1087	SEQ ID No. 583	SEQ ID No. 584	SEQ ID No. 1088	SEQ ID No. 1090	SEQ ID No. 3112	SEQ ID No. 3273	SEQ ID No. 2761	SEQ ID No. 3272	D No.	Ö. So.	ID No.	Ö No	SEQ ID No. 3446	SEQ ID No. 2760

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35 (2 IV) 08	102.1	Oction	a.	34385	3/093		ı	%Q6	
SEQ ID No. 75	103.1	Contig50	۵	37068	38339	+		%86	
SEQ ID No. 83	104.1	Contig50	a.	38294	39469	+		%86	
SEQ ID No. 103	107.1	Contig50	۵	39535	42687		•	%86	
SEQ ID No. 117	109.1	Contig50	a	42543	44045	ı	•	94%	,
SEQ ID No. 133	111.2	Contig50	Ω	43985	45556	•	•	%96	,
SEQ ID No. 851	227.2	Contig50	a	45523	46464		,	93%	ı
SEQ ID No. 859	228.1	Contig50	Q.	46477	46656		•	%26	+
SEQ ID No. 892	234.2	Contig50	a .	46705	48930	1	ı	%66	,
SEQ ID No. 680	1961.3	Contig50	d.	49060	49950		ı	%96	+
SEQ ID No. 390	152.3	Contig50	a	50084	50461	+		%86	ı
SEQ ID No. 398	153.2	Contig50	ď	50462	51799	+	1	%66	·
SEQ ID No. 405	154.1	Contig50	۵	51796	52764	+	1	%96	•
SEQ ID No. 424	157.2	Contig50	۵	52765	55980	+	1	%86	1
SEQ ID No. 3107	594.1	Contig50	۵	55956	56429	+		%96	+
SEQ ID No. 3105	593.2	Contig50	٤	56568	56918		1	%96	+
SEQ ID No. 3102	592.3	Contig50	۵	57113	58795	•	ı	94%	,
SEQ ID No. 504	169.1	Contig50	Ε	58808	59020	,	+		+
SEQ ID No. 497	168.1	Contig50	۵	59115	29687	,	1		
SEQ ID No. 486	166.2	Contig50	۵	59537	62800		1	95%	
SEQ ID No. 464	163.1	Contig50	a	62978	64276	,	+		•
SEQ ID No. 447	160.1	Contig50	ε	64504	64863		+		
SEQ ID No. 438	159.1	Contig50	۵	65005	65316		t	%68	,
SEQ ID No. 430	158.2	Contig50	۵	65304	65987		ı	%86	
SEQ ID No. 679	1960.2	Contig50	Q.	65972	66484	+	•	92%	+
SEQ ID No. 462	1628.3	Contig50	E	66550	67515	+	1	92%	•
SEQ ID No. 463	1629.1	Contig50	٤	67365	68138	+	1	95%	•
SEQ ID No. 465	1631.3	Config50	٤	68126	70117		ı	%26	1
SEQ ID No. 1091	2637.1	Contig50	۵	70315	70731		•	100%	+
SEQ ID No. 3325	818.2	Contig50	٤	70763	71647		ı	94%	+
SEQ ID No. 3326	819.2	Contig50	E	71690	73171	3	ı	100%	•
SEQ ID No. 411	1550.2	Contig50	E	73149	73937	,	,	%26	+
SEQ ID No. 410	1549.1	Contig50	٤	73898	74191	+	1	100%	,
D No.	1548.2	Contig50	٤	74170	74871			%66	r
D No.	1425.2	Contig50	٤	74865	75158		t	%16	. •
SEQ ID No. 335	1424.2	Contig50	٤	75133	75561		1	%96	1

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75552	77314	78822	79435	79937	81725	82659	83861	84690	84994	85582	85971	86294	87124	87816	88330	90968	90587	91012	91731	93538	94413	95685	95702	96880	98100	98526	99459	100596	101324	102287	102978	103508	104476	105120	70000
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1423.2	2639.1	3226.2	2024.2	148.2	150.2	5473.1	151.2	3228.1	3230.1	3231.1	3232.1	3233.1	3234.3	5474.2	875.5	876.1	879.2	5615.1	5920.2	704.4	705.1	706.2	707.2	1992.2	3472.1	3473.1	3476.1	3477.2	3633.3	3634.1	3636.1	3637.1	3638.2	3640.2	1017.2
SEQ ID No. 334	SEQ ID No. 1092	SEQ ID No. 1470	SEQ ID No. 708	SEQ ID No. 363	SEQ ID No. 376	SEQ ID No. 2902	SEQ ID No. 382	SEQ ID No. 1471	SEQ ID No. 1473	SEQ ID No. 1474	SEQ ID No. 1475	SEQ ID No. 1476	SEQ ID No. 1477	SEQ ID No. 2903			₽	₽	₽	SEQ ID No. 3245	SEQ ID No. 3246	SEQ ID No. 3247	SEQ ID No. 3248	SEO ID No. 693		SEQ ID No. 1638		SEQ ID No. 1640	SEQ ID No. 1753	SEQ ID No. 1754	SEQ ID No. 1755			SEQ ID No. 1759	SEO ID No 673

107274 109404 109404 110134 1110392 112805 115604 11968 11968 11968 11968 12362 12797 12797 12797 1368 1318 1318 1340 1340 1353 1363 1363 1363 137 140 141 141 141 141 141 141 115459 116708 118435 119212 119787 121146 122748 125624 125624 126853 128135 130769 131938 134740 135659 135939 13666 138666 106981 107472 108772 109367 110241 111723 Contig50
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SEQ ID No. 2484
SEQ ID No. 2487

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SEQ ID No. 1015	2510.2	Contig50	٥	144653	146515	•	ı	%26	•
SEQ ID No. 1013	2509.3	Contig50	. മ	146573	147010		ı	%86	+
SEQ ID No. 1411	3129.2	Contig50	م	147083	148138	,	1	%66	
SEQ ID No. 1412	3130.1	Contig50	۵	148182	149018		1	%26	1
SEQ ID No. 1413	3131.1	Contig50	۵	149155	149499	+	•	100%	1
ģ	949.3	Contig50	Ф	149500	151377	+	•	+ %66	1
D No.	2148.1	Contig50	ď	151378	152307	1		+ 4 4 4 4	
SEQ ID No. 779	2147.1	Contig50	Ε	152485	152835		•	100%	•
SEQ ID No. 778	2146.2	Contig50	Ε	152863	154005		1	%86	
SEQ ID No. 1414	3132.1	Contig50	Ф	154060	154446			. 95%	+
D No.	3133.2	Contig50	a	154385	155191	1	1	%26	+
№	5269.1	Contig50	ď	155067	155423		1	94%	+
ID No.	5270.2	Contig50	Ф	155164	155745		ı	95%	+
D No.	4372.3	Contig50	۵	155848	157050	+	1	%86	ı
ID No.	4371.1	Contig50	٤	157168	158067		•	%66	•
ID No.	4370.1	Contig50	a	158463	159020	1	•	%86	+
Š	4369.1	Contig50	م	159079	159750		1	88%	. •
Б В	4367.1	Contig50	а	159936	160592	ı	•	82%	+
Ω	4366.1	Contig50	Ε	160879	161298	•	•	91%	i
Θ. No	4365.4	Contig50	E	161394	161951		1	%98	1
D No	5602.3	Contig50	d	162005	162481	1		%26	•
<u>8</u>	808.4	Contig50	a	162640	165525	1		78%	+.
₽	807.2	Contig50	a.	165692	167020		,	93%	. 1
D No.	806.2	Contig50	٤	167060	167803	ı	1	%26	ı
Ю В	4082.2	Contig50	E	167817	169130	1		95%	•
Ю В	1726.4	Contig50	Ε	169200	170873			93%	J
Ю В	5484.4	Contig50	Ε	170954	172321		,	94%	+
ID No.	4567.2	Contig50	<u>a</u>	172530	174488		•	91%	•
	4566.1	Contig50	٤	174558	175169	,	1	%96	,
D No	5901.3	Contig50	Ε	175331	176827			82%	
SEQ ID No. 3093	5900.3	Contig50	a	176579	176953		•	84%	+
SEQ ID No. 2444	4663.3	Contig50	Ε	177123	178034	+		%26	,
Ю Мо	4662.1	Contig50	Ε	178485	178745		+		+
Ю В	4660.2	Contig50	Ε	179180	179395		+		+
D No.	4659.2	Contig50	d	179570	179950		ı	93%	+
SEQ ID No. 2440	4658.2	Contig50	۵	180138	180596		ı	%96	ı

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181254	183482	184141	184968	185581	186055	187947	188745	189430	753	2896	4694	5275	5569	6735	7809	8312	8610	8743	9696	10940	11305	12434	13476	14061	15003	15778	16097	16500	16967	18522	19188	19598	19821	19837	20172
180604	181581	183848	184198	185006	185774	186391	188128	188912	46	1154	3624	4604	5276	5479	2069	7842	8263	8510	8710	9882	11030	11064	13063	13669	14167	15275	15834	16060	16809	17140	18871	19419	19507	19664	19993
۵	<u>a</u>	۵	Ε	E	٥	Q.	E	Ω.	۵	E	۵	<u>a</u>	O.	a	Ω.	٤	ď	а	ď	۵	۵	٤	٤	٤	٤	E	Ε	ε	Ε	Q.	a	Ε	۵	Ε	۵
Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig50	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51
4657.1	4656.2	6064.1	5344.2	5342.2	5341.1	5340.1	4984.2	6062.1	3314.2	3316.1	2057.3	977.3	976.3	974.2	2056.1	1270.2	1271.3	5983.1	1272.4	2054.2	5467.1	3319.1	4962.2	4961.1	4960.2	4943.2	4942.1	4941.2	2141.2	748.3	749.1	750.2	751.2	752.2	6061.1
D No O	SEQ ID No. 2438	SEQ ID No. 3151	SEQ ID No. 2842	SEQ ID No. 2841	SEQ ID No. 2840	SEQ ID No. 2839	SEQ ID No. 2657	SEQ ID No. 3150	ID No.			№ №	ID No.		D So	SEQ ID No. 234	SEQ ID No. 235	D No	SEQ ID No. 236	D No	Б В	Ю В	SEQ ID No. 2645	Ö So	Б В	è S S	<u>8</u> Q	Ö S O	D No.	Ю В	SEQ ID No. 3275		Ö No.		SEQ ID No. 3149

96% 99% 99% 98% 83% 21584 22934 23767 24621 24621 25828 27000 29141 30647 33643 34675 35431 43565 43565 43565 43565 43565 43565 50593 50593 50657 51767 55263 20064 21993 21993 22973 22973 22973 22903 30907 31862 33284 34700 33572 34700 34700 34700 34700 34700 3572 36572 44028 49013 5674 5673 5673 Contig51
Contig51 2143.5 1182.3 1180.2 1180.2 1180.2 1179.1 1179.1 2712.1 2717.1 2717.1 1324.4 1385.2 2724.3 2724.3 2724.3 2724.3 3442.3 3442.3 3442.3 3442.3 3445.3 SEQ ID No. 776
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183347 183853 183853 185502 185887 186597 186597 190029 190493 191123 191123 191123 191123 195566 195566 195566 1955845 196133 196133 198343 19837 200154 200154 2004547 205471 206756 207652 208368 210298 210398 191412 194051 195204 195579 195800 196361 199397 199903 200543 202374 203513 204548 205599 190797 191116 185959 186827 187570 188799 189586 190023 E E E O E E E E E E E E E E E O O O E O O O E E E O E E E E O E Contig51 123.2 124.1 125.2 2022.1 2020.1 2020.1 2020.1 2020.1 2020.1 2020.1 2020.1 4856.2 4856.2 4856.2 4856.2 4856.2 4856.2 4856.2 4856.2 4856.2 4856.2 3516.2 3517.1 3517.1 3509.2 3509.1 3509.1 3509.1 3509.1 3509.1 3509.1 3509.1 3509.1 SEQ ID No. 216
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213997	214705	216364	217278	218064	219146	220517	221437	222089	222642	223928	224949	226279	228321	224	358	1145	1785	2453	2762	3519	4989	6110	6424	6950	7735	9080	8966	10775	11494	12546	13507	15502	16920	18783	10206
Q	Ε	a	Ω.	Ω	α.	Ε	Ε	Ε	Ε	a.	Ε	Ε	٤	а	Q.	٤	E	E	Ε	a	Ε	Ω.	a	۵	۵	۵	۵	۵	۵	۵	٥	۵	ď	۵	
Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig51	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contin52									
1521.2	4415.3	4325.3	4326.1	4327.1	4328.1	4740.2	4739.1	4737.1	4736.2	4734.2	5440.1	64.3	65.1	5556.1	5555.2	6052.1	5201.2	5202.2	5204.2	4444.2	4442.1	4441.1	4440.1	1863.2	1862.2	801.3	802.3	803.3	5206.1	3142.2	1532.3	1533.1	1246.2	1245.1	1244 2
ID No.	Ö N O	SEQ ID No. 2205	SEQ ID No. 2206	SEQ ID No. 2207	SEQ ID No. 2208	SEQ ID No. 2499	SEQ ID No. 2497	SEQ ID No. 2496	SEQ ID No. 2495	SEQ ID No. 2494	SEQ ID No. 2887	SEQ ID No. 3202	SEQ ID No. 3209	SEQ ID No. 2951	SEQ ID No. 2950	SEQ ID No. 3147	SEQ ID No. 2778	SEQ ID No. 2779	SEQ ID No. 2780	SEQ ID No. 2295	SEQ ID No. 2294	SEQ ID No. 2293	SEQ ID No. 2292	Ω	ID No.	Ω		Ю В	© So.	₽	SEQ ID No. 401			SEQ ID No. 219	SEO ID No 218

SEQ ID No. 329	1417.1	Contig52	۵	20421	20891		,	100%		
SEQ ID No. 330	1418.3	Contig52	· E	20994	22823	+	•	%06		
SEQ ID No. 3146	6049.1	Contig52	Ε	22855	23631	ı	+			
D So.	2954.3	Contig52	Ε	23498	24508	ı	1	%66	٠	
ID No.	601.4	Contig52	۵	24684	25916	,	+			,
Ω So	600.3	Contig52	۵	25859	26539		+			
Θ Ω	598.2	Contig52	۵	26619	26939	+	1	100%		
Ю В	597.2	Contig52	۵	26966	27346	+	ı	%86		
№ №	596.3	Contig52	Ε	27403	29127		1	%66	+	
0 №	2955.2	Contig52	٩	29162	29581		•	%66		+
№ №	2958.2	Contig52	۵	29575	30342		1	%86		
ID No.	2959.1	Contig52	Q.	30346	31104	+	1	%26		
D No	2960.1	Contig52	۵	31112	32215		1	%86		
D No.	2961.1	Contig52	٤	32318	33571	,	1	%86		
D No.	2962.1	Contig52	Q.	33682	34458		ı	%86		
ID No.	2495.2	Contig52	۵	34434	34982	,		%86		
D No	2493.2	Contig52	٥	34913	35578		ı	%86		
№ №	2492.2	Contig52	۵	35616	35948	+	1	%56	·	+
ID No.	938.3	Contig52	٤	35989	37614		1	%06		
D No	937.3	Contig52	٤	37875	39305	+	ı	%66		
В В	1589.3	Contig52	٤	39287	39994	1		%66		
Ю В	1590.1	Contig52	٤	40025	40582	+	•	%86		
D No.	1591.4	Contig52	٤	40813	42264		•	%86		
Ö. Se	2963.2	Contig52	٤	42366	43133		•	%86		
D No.	2964.1	Contig52	٤	43134	43991		ı	%86		
D No.	2965.1	Contig52	a.	44163	44477	1	1	100%	•	+
D No.	2966.1	Contig52	Ε	44480	45376	1	•	%26		_
≥ Ω	2967.1	Contig52	Ε	45421	45705	,	•	100%	•	_
Б В	2968.3	Contig52	a	45890	48583		•	%92	•	
D No.	5455.1	Contig52	Ε	48635	49105			%86		
D No.	2156.3	Contig52	٤	49102	50409		ı	%86	•	
Θ S O	1740.3	Contig52	a	50644	51432		•	100%	•	
В В	1739.2	Contig52	d	51433	52056		1	%66	•	
Ö S S	1738.3	Contig52	a	52040	53257	r	ı	%66	+	
S O	1547.2	Contig52	ď	53250	54101		•	%66	•	
SEO ID No. 407	1545.3	Contig52	Q.	54083	55753	1	•	%66	+	

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55754	57282	59352	59770	61258	62267	63680	64855	67104	68261	69468	71343	72814	73211	75741	77475	78868	79583	80548	81928	82091	83467	84536	84956	86644	87008	88842	89686	90941	69606	91747	92270	92893	93920	94512	97049
ο.	Ω.	Q .	٤	٤	۵	a	E	٤	٤	٤	٤	۵	٤	٤	٤	Ε	Ε	Ε	Ε	۵	٤	٤	٤	٤	٤	a	Q.	٤	a	a	٤	٤	٤	Ε	٤
Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52
3311.1	2235.2	3312.1	5534.2	4197.2	5535.1	1448.2	1449.3	4195.3	5100.2	3378.2	3376.1	366.2	368.2	3374.1	982.3	983.1	985.2	3371.2	4813.2	3368.2	4814.2	4815.2	130.4	129.3	126.2	465.2	464.2	463.2	462.2	460.2	459.3	458.3	2943.1	2942.1	2941.1
SEQ ID No. 1531	SEQ ID No. 831	SEQ ID No. 1532	SEQ ID No. 2937	SEQ ID No. 2121	SEQ ID No. 2938	ID No.	D No.	SEQ ID No. 2120	SEQ ID No. 2734	SEQ ID No. 1578	SEQ ID No. 1577	SEQ ID No. 1773	SEQ ID No. 1788	SEQ ID No. 1576	SEQ ID No. 3441	SEQ ID No. 3442	SEQ ID No. 3443	SEQ ID No. 1575	SEQ ID No. 2553	SEQ ID No. 1573	SEQ ID No. 2554	SEQ ID No. 2555	SEQ ID No. 255	SEQ ID No. 249	SEQ ID No. 227	SEQ ID No. 2433	SEQ ID No. 2425	SEQ ID No. 2420	SEQ ID No. 2412	SEQ ID No. 2400	SEQ ID No. 2393	SEQ ID No. 2386	SEQ ID No. 1279	SEQ ID No. 1278	SEQ ID No. 1277

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98600 99826	101356	103845	105045	106336	109404	110751	113277	114581	115254	115556	116104	116444	117759	119229	119802	120579	121508	123297	125370	125799	127072	128015	128793	130032	131921	133076	133824	134192	135727	135446	137590	139124	140526	141347
98412 98687	99827	101440	103846	105158	106495	109609	110890	113583	114538	115152	115526	115998	116524	117790	119230	119803	120600	121603	123484	125482	125894	127035	128218	128905	130020	132081	133339	133818	134261	135288	135914	137514	139087	140406
<u>a</u> a	o.	Q.	Q	Ε	Ε	۵	٤	Ε	Ε	Ε	Ε	Ε	ε	۵	۵	۵	۵	۵	۵	Ε	٤	Ε	۵	Ε	٤	۵	E	٤	Ε	a	Ω	۵	۵	۵
Contig52 Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52	Contig52
5944.1 1215.3	1220.3	2103.2	1638.2	1639.4	1635.4	5481.1	4665.2	4288.2	4289.2	4291.1	770.3	771.4	772.4	774.3	775.2	4293.2	5482.2	3909.3	3911.2	2382.2	2381.2	2377.2	2375.3	3913.2	3914.3	2239.3	2240.2	2241.2	4384.2	6044.1	4385.1	4386.2	4867.2	4868.2
SEQ ID No. 3108 SEQ ID No. 198	SEQ ID No. 200	Ω		\Box	SEQ ID No. 468	SEQ ID No. 2909	SEQ ID No. 2445	SEQ ID No. 2178	SEQ ID No: 2179	SEQ ID No. 2181	SEQ ID No. 3293	SEQ ID No. 3294	SEQ ID No. 3295	SEQ ID No. 3296	SEQ ID No. 3297	SEQ ID No. 2182	SEQ ID No. 2910	SEQ ID No. 1920	SEQ ID No. 1921	D No.	SEQ ID No. 924		SEQ ID No. 921		SEQ ID No. 1923	SEQ ID No. 832	SEQ ID No. 833	SEQ ID No. 834	SEQ ID No. 2253	SEQ ID No. 3145	SEQ ID No. 2254	SEQ ID No. 2255	SEQ ID No. 2585	SEQ ID No. 2586

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141335 141660	142688	143354	143907	145418	145945	147294	147706	147984	149012	149995	150975	151758	152038	153313	154287	154943	155962	156342	157275	158419	160244	161680	163009	164484	165688	166701	167624	168334	168692	170259	171843	173322	174266	176847
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4869.3 5549.2	5548.1	5546.1	4874.2	4875.1	4877.2	4629.2	4628.2	4627.2	4626.1	4625.2	4803.2	4804.1	4805.1	4806.4	6042.1	4958.3	4957.2	1346.3	1345.3	3517.3	1450.2	3519.1	3520.1	3521.2	3522.1	3524.1	3525.1	3527.1	4228.2	4227.1	1059.2	1060.2	4226.2	2395.3
Б No.	SEQ ID No. 2944	SEQ ID No. 2943	SEQ ID No. 2590	D No.	SEQ ID No. 2592	SEQ ID No. 2419	SEQ ID No. 2418	SEQ ID No. 2417	SEQ ID No. 2416	ġ	SEQ ID No. 2545	SEQ ID No. 2546		SEQ ID No. 2548	SEQ ID No. 3144	SEQ ID No. 2641		SEQ ID No. 282	SEQ ID No. 281		SEQ ID No. 347	SEQ ID No. 1671	ω So	ID No.	0 №	D No	Ю В	\Box	ID No.		Ω So	Ö. Se	D No.	SEQ ID No. 932

190959 191497 193376 194182 195009 1956262 197824 198467 199365 199801 200760 201863 202480 203940 203940 205314 205314 206050 2011148 179017 179285 179977 181719 182123 183339 184718 185408 185408 188973 189461 190114 195444 196424 197784 198472 199322 199795 201015 202699 203934 204682 205289 205289 207457 210147 188964 189554 190081 191093 191571 193397 194158 195010 85529 Contig52
Contig52 Contig52 Contig52 1611.5 6039.1 6039.1 6039.1 4489.3 4489.3 4488.1 4487.1 4480.2 4800.2 5587.3 4800.2 5103.4 5104.3 7712.2 3712.2 3710.1 3709.1 3703.1 3703.1 3703.1 3703.1 3703.1 3703.1 SEQ ID No. 3143
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SEQ ID No. 2678	5021.3	Contig52	. ∈	229852	230601			97.70	+
SEQ ID No. 2676	5019.1	Contig52	Ε	230836	231474	,	ı	%00 %00	
SEQ ID No. 2675	5018.2	Contig52	٤	231413	231898			%66 0000	1
SEQ ID No. 2464	4696.3	Contig52	Ε	231883	233508		1 1	%0%	+
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SEQ ID No. 444	1598.6	Contig52	<u> </u>	236937	237020	ı		%86,	•
SEQ ID No. 443	1595.3	Contig52	2 0	237930	230125		•	%00L	•
SEQ ID No. 442	1594.2	Contig52		239053	230272	i		%86	ı
SEQ ID No. 989	2472.2	Contig52		239763	240410		t	%/6	
SEQ ID No. 990	2473.1	Contig52	LΩ	240545	240859		! 1	96%	ı
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S S	4376.3	Contig52	۵	246162	246848	,	ı	90.70	+
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S O	6038.1	Contig52	Ω.	247180	247419		,	%96 	+ -
Ö. So	971.2	Contig52	Ε	248160	249020			%86	+ ·
SEQ ID No. 3434	972.3	Contig52	۵	249094	250731		ı	%96 %90	+
Ω	4891.2	Contig52	E	250785	251615		ı	%00 %00	•
Ö N	4889.1	Contig52	٤	251793	252218	,	ı	400%	•
SEQ ID No. 2601	4888.1	Contig52	٤	252280	252804	,	•	%96 %96	,

SEQ ID No. 3341	843.2	Contig52	ε	295435	296004	ı	,	%66	
SEQ ID No. 3342	844.2	Contig52	٤	296047	296436	1	1	%66	٠,
SEQ ID No. 3343	845.4	Contig52	٤	296556	297314		•	%86	1
SEQ ID No. 1002	249.4	Contig52	a	297549	299282	1	ı	%86	
SEQ ID No. 3071	5848.1	Contig52	٤	298199	298495		•	%96	+
SEQ ID No. 1008	250.1	Contig52	σ.	299279	300508	•	•	%66	
ID No.	251.2	Contig52	Ф	300542	301417	ı	,	%66	+
№	252.2	Contig52	E	301431	302669		1	%86	
Ю No	3898.2	Contig52	Ε	302744	303340		1	%86	+
<u>Б</u>	3899.2	Contig52	Ε	303353	303835		,	%86	+
© No	3901.2	Contig52	<u>α</u>	303849	304757		t	%66	
SEQ ID No. 1917	3902.3	Contig52	Ε	304877	306370		í	%86	
ID No.	3857.3	Contig52	Ε	306327	307457	+	ı	%86	,
В	3855.1	Contig52	٤	307661	308464		•	%86	
ID No.	1398.2	Contig52	٤	308532	310538		•	%26	
SEQ ID No. 316	1399.3	Contig52	Ε	310525	312153		1	%66	1
№	3853.2	Contig52	٤	312450	313760		1	%86	ı
ID No.	3851.1	Contig52	d.	314121	314300		+		
D So.	3850.1	Contig52	E	314297	314740	•	ı	%26	
№ №	3849.1	Contig52	Ε	314722	315453		•	%66	ı
	4863.4	Contig52	۵	315679	317031		1	%66	1
Б В	4864.2	Contig52	d	317013	317423		+		ı
ID No.	5453.2	Contig53	Ε	7	1057	+	,	%66	ı
ID No.	3244.2	Contig53	۵	1210	1977	,	1	%66	ı
SEQ ID No. 3187	618.3	Contig53	۵	2045	2461		ı	100%	ı
D So	619.1	Contig53	۵	2371	3357		ı	%26	•
D No	621.5	Contig53	ď	3335	4654		ı	%66	1
S S	687.5	Contig53	ď	4655	5932	+	ı	%86	•
Ю В	688.1	Contig53	۵	5916	6770		ı	%66	1
<u>.</u> №	3243.1	Contig53	Ε	0889	7188		ı	%66	+
Ю	3242.3	Contig53	Ε	7361	10084	ď	1	%86	+
SEQ ID No. 2989	5633.2	Contig53	E	10360	11202		ı	%86	1
N S	5634.2	Contig53	E	11203	11337	+	+		
<u>.</u> <u>S</u>	5636.1	Contig53	٤	11490	11912		•	%66	+
ID No.	5637.1	Contig53	α.	12003	12689	1	,	%66	ı
SEQ ID No. 2516	4766.2	Contig53	ď	12665	13120	ı	•	100%	

SEQ ID No. 2517	4767.2	Contia53	۵	13062	13379			100%	•
SEQ ID No. 2518	4768.1	Contig53	. E	13467	13820		•	%66 	ı
SEQ ID No. 2519	4769.1	Contig53	а	14000	14419		•	100%	٠,
SEQ ID No. 2520	4770.1	Contig53	a	14512	14700	ı	+		1
SEQ ID No. 2521	4771.2	Contig53	ď	14837	15913			%96	1
SEQ ID No. 2993	5638.2	Contig53	Ε	16224	16634	+	,	100%	+
SEQ ID No. 2854	5377.2	Contig53	۵	16834	17466		,	95%	+
SEQ ID No. 2853	5376.2	Contig53	Ε	17524	21840			%66	+
SEQ ID No. 1375	3077.1	Contig53	Ε	21958	22719		ı	%26	ı
SEQ ID No. 1376	3078.2	Contig53	ε	22682	23938		,	· %66	ı
SEQ ID No. 1377	3080.2	Contig53	ε	23895	25349			100%	ı
SEQ ID No. 1378	3081.2	Contig53	Ε	25561	26097			%86	+
SEQ ID No. 1379	3082.1	Contig53	Ε	26262	26786			%86	
SEQ ID No. 1380	3083.1	Contig53	Ε	26732	27307	+		%26	1
SEQ ID No. 840	2251.3	Contig53	Ε	27542	28039			95%	+
SEQ ID No. 1381	3085.2	Contig53	d	28077	30950	,	•	+ %86	
SEQ ID No. 1382	3086.1	Contig53	۵	31106	31714	+	•	100%	•
SEQ ID No. 1383	3087.1	Contig53	۵	31707	32744		•	%66	1
SEQ ID No. 1384	3088.1	Contig53	Ε	32771	33550		ı	100%	ı
SEQ ID No. 2594	488.2	Contig53	Ε	33534	34451	ı	ı	%66	1
SEQ ID No. 2603	489.1	Contig53	Ε	34530	34715	1	+		+
SEQ ID No. 2613	491.1	Contig53	٤	34679	35737	ı	•	%26	+
SEQ ID No. 2620	492.4	Contig53	Ε	35824	36363		•	%86	1
SEQ ID No. 2852	5373.1	Contig53	E	36255	36521		+		+
Š	493.4	Contig53	Ε	36559	37296		•	%96	
SEQ ID No. 1386	3091.1	Contig53	٤	37453	37764		,	100%	+
SEQ ID No. 3075	5867.1	Contig53	۵	37577	37750		•	94%	+
Ω	1705.2	Contig53	ė	37874	38746	1	•	%86	+
Ω	1706.2	Contig53	٤	38847	39317	ı		%86	ı
Ö S O	1707.2	Contig53	Ε	39318	39686		ı	100%	1
D No.	3093.1	Contig53	٤	39710	40480	1		%86	•
<u>0</u>	3095.1	Contig53	٤	40462	41001	ı		%86	1
D No.	3096.1	Contig53	٤	40977	41246	1	•	100%	ı
SEQ ID No. 1390	3097.3	Contig53	٤	41324	42712	ı	•	%66	•
EQ ID No.	3960.2	Contig53	٤	42957	43676		•	93%	+
SEQ ID No. 1945	3959.2	Contig53	۵	43842	45401		ı	%16	

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06230	43764	4/095	47919	48480	48942	49907	50175	50884	51398	52389	53322	54335	55250	55808	57129	57980	59751	60784	61569	62888	63083	64558	64881	65548	66654	66802	67100	67640	68083	69533	70542	71489	72395	72795	73488	74253
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7.42.3	742.0	741.2	740.2	2213.2	3958.1	3957.1	3956.2	5524.1	5523.2	2400.2	2401.2	1476.2	1475.2	1989.2	1990.1	241.2	240.1	239.1	238.1	236.1	235.2	2969.1	2970.2	916.3	917.3	918.3	919.1	920.2	4655.3	1353.3	1354.2	4653.2	4652.2	4432.2	4433.1	4434.1
SEO ID No. 3270			SEQ ID No. 3268	SEQ ID No. 819	SEQ ID No. 1944	SEQ ID No. 1943	SEQ ID No. 1942	SEQ ID No. 2929	SEQ ID No. 2928	SEQ ID No. 935	SEQ ID No. 936	SEQ ID No. 361	SEQ ID No. 360	SEQ ID No. 690	SEQ ID No. 692	SEQ ID No. 942	SEQ ID No. 934	SEQ ID No. 929	SEQ ID No. 923	SEO ID No. 908	SEQ ID No. 898	SEQ ID No. 1299	SEQ ID No. 1301	SEQ ID No. 3392	SEQ ID No. 3393			SEQ ID No. 3396	ġ	ġ	ġ	SEQ ID No. 2436	SEQ ID No. 2435	В	Ω So	SEQ ID No. 2289

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75252	78146	79261	79662	79932	80346	80802	81579	82655	85470	86027	86339	86938	87390	88067	89749	91234	92027	92655	93248	93919	94806	95892	97055	98314	101235	102305	103921	104792	106148	106644	107241	109621	111272	112570	113522
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Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53	Contig53
4435.4	451.3	450.1	448.1	447.2	444.2	443.3	3562.3	3561.2	874.2	873.2	872.2	870.3	869.2	668.4	667.3	3322.2	3323.1	3324.1	434.3	433.1	432.1	431.1	1071.3	1072.4	3326.2	5238.1	940.4	941.3	943.3	944.3	3928.1	3929.2	3814.1	969.2	967.2
SEQ ID No. 2290		S : ⊡	S S S		D No.	SEQ ID No. 2285	SEQ ID No. 1703	SEQ ID No. 1702				SEQ ID No. 3359	SEQ ID No. 3358	SEQ ID No. 3222	SEQ ID No. 3221	SEQ ID No. 1536	SEQ ID No. 1537	SEQ ID No. 1538	SEQ ID No. 2217	SEQ ID No. 2209	SEQ ID No. 2200	SEQ ID No. 2194	SEQ ID No. 104	SEQ ID No. 105	SEQ ID No. 1539	SEQ ID No. 2794	SEQ ID No. 3411	9.	9.	<u>و</u> :		D No	و 0	S S S S	SEQ ID No. 3430

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151252	151701	152475	152886	154337	154705	155302	156234	157090	158359	159194	159545	159996	160720	161325	161789	162784	163191	164165	164821	165348	166781	167095	167507	169298	170063	172891	174213	175448	176752		177979	177979 178463	177979 178463 179471	177979 178463 179471 180979	177979 178463 179471 180979 182641
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730.1	729.1	728.2	3024.1	3023.1	3022.1	1268.2	1267.3	1265.5	3018.1	3017.1	3016.1	3015.1	3014.1	3013.2	3011.2	5288.1	5289.2	5312.2	5313.1	4897.2	4895.1	4894.1	1945.4	1946.2	1157.4	3468.2	1559.2	1557.3	2356.2		1786.2	1786.2 1784.2	1786.2 1784.2 1782.3	1786.2 1784.2 1782.3 5316.1	1786.2 1784.2 1782.3 5316.1 5123.3
SEQ ID No. 3262	SEQ ID No. 3261	₽	₽	₽	SEQ ID No. 1339	SEQ ID No. 233	SEQ ID No. 232	SEQ ID No. 231	SEQ ID No. 1338	SEQ ID No. 1337	SEQ ID No. 1336	SEQ ID No. 1335	SEQ ID No. 1334	SEQ ID No. 1333	SEQ ID No. 1332	SEQ ID No. 2814	SEQ ID No. 2815	SEQ ID No. 2824	SEQ ID No. 2825	SEQ ID No. 2607	SEQ ID No. 2606	SEQ ID No. 2605	SEQ ID No. 671	SEQ ID No. 672	SEQ ID No. 161	SEQ ID No. 1635	SEQ ID No. 416	SEQ ID No. 415	SEQ ID No. 905		SEQ ID No. 571	\Box	SEQ ID No. 571 SEQ ID No. 570 SEQ ID No. 569		SEQ ID No. 571 SEQ ID No. 570 SEQ ID No. 569 SEQ ID No. 2826 SEQ ID No. 2743

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183405	184332	184907	185478	187318	189743	190543	191202	192709	193404	194228	194570	195867	196513	197390	198224	198889	199470	200568	201197	202045	203633	205210	205742	205998	206346	206934	209993	210303	210846	212435	213767	214284	215547	216391	217448
Ε	Ε	۵	Ω.	E	Ε	Ε	Ε	ε	ε	E	۵	Ε	۵	Q.	۵	۵	Ε	۵	م	٤	a	E	Ε	۵	۵	a	Ω.	٤	٤	٤	۵	٤	Ε	<u>α</u>	a.
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5127.4	3982.2	3980.1	3979.2	3977.2	1494.2	1495.1	1497.2	5317.2	1606.2	1607.2	1608.2	4597.1	4598.2	4729.3	4727.1	4726.1	4725.2	4724.2	4721.2	4878.2	2859.2	2858.1	81.3	80.3	2743.2	2745.1	692.1	693.1	694.3	2747.1	2748.1	2749.4	1760.4	1759.4	3591.2
SEQ ID No. 2745	SEQ ID No. 1964	SEQ ID No. 1963	SEQ ID No. 1961	SEQ ID No. 1960	SEQ ID No. 372	SEQ ID No. 373	SEQ ID No. 374	SEQ ID No. 2827	SEQ ID No. 451	SEQ ID No. 452	SEQ ID No. 453	SEQ ID No. 2396	SEQ ID No. 2397	ġ	SEQ ID No. 2489	SEQ ID No. 2488	SEQ ID No. 2487	SEQ ID No. 2486	SEQ ID No. 2485	SEQ ID No. 2593	SEQ ID No. 1228	SEQ ID No. 1227	SEQ ID No. 3321	9	Ω	SEQ ID No. 1160	₽	SEQ ID No. 3236	SEQ ID No. 3237	SEQ ID No. 1161	SEQ ID No. 1162	SEQ ID No. 1163	SEQ ID No. 552	SEQ ID No. 550	SEQ ID No. 1720

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218007	219507	220002	221578	221926	222331	223164	223351	224214	225046	227254	227495	227848	228193	230355	230827	232262	233014	234284	235534	236025	236327	237431	238592	239239	241173	242561	243808	244324	244779	245702	246861	248091	248604	249860
۵ ۵	Ε	Ε	Ε	۵	Ε	Ω.	۵	ε	۵	۵	۵	ε	ε	۵	Ω	Q.	ο.	۵	۵	a	a.	٤	Ε	Ε	۵	٤	Ε	۵	Ε	Q.	O.	٤	ď	Ф
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3592.1 3593.1	3594.1	2421.2	3595.1	3596.1	3597.3	3598.3	3599.4	5741.1	2657.3	2655.1	2654.1	2653.1	482.2	481.1	480.3	1951.4	1950.2	342.2	343.1	344.1	346.2	347.1	349.2	2652.2	2650.1	2521.4	2522.1	2523.1	2524.2	2649.1	650.3	651.2	652.1	2648.2
SEQ ID No. 1721 SEQ ID No. 1722	ID No.	SEQ ID No. 950	SEQ ID No. 1724	SEQ ID No. 1725	SEQ ID No. 1726	D No.	SEQ ID No. 1728	SEQ ID No. 3039	ω Ω	SEQ ID No. 1105	SEQ ID No. 1104	SEQ ID No. 1103	SEQ ID No. 2559	SEQ ID No. 2550	SEQ ID No. 2541	SEQ ID No. 677	SEQ ID No. 676		SEQ ID No. 1612	SEQ ID No. 1616	SEQ ID No. 1630	SEQ ID No. 1636	SEQ ID No. 1649	ID No.	SEQ ID No. 1101	Б В	ID No.	SEQ ID No. 1023	SEQ ID No. 1024	SEQ ID No. 1100	SEQ ID No. 3210		D No.	SEQ ID No. 1099

266431 267495 268047 268976 270013 271638 273283 273981 275271 275571 276554 277948 277852 280065 281205 281789 262400 265922 261939 Contig53
Contig53 2647.2 2646.1 2645.1 1288.2 1915.3 1915.3 1915.3 1915.3 1913.2 2643.1 1926.2 1926.2 1926.2 1926.2 1926.2 3859.2 3859.2 2688.2 2688.2 3860.2 3860.2 3860.2 3860.2 3860.2 3860.2 3860.2 1926.2 1926.2 1926.2 1926.2 1926.2 1926.2 1926.3 1926.2 1926.3 1926.2 1926.3 1926.2 1926.3 19 SEQ ID No. 1096
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7698	9281	10058	10008	11108	11802	12803	13131	13347	13646	13937	16404	17111	17664	18708	19421	20170	21258	22264	24158	25053	26048	28868	29266	29522	30263	31076	31582	33023	33345	34066	34408	34645	35802	36853	37802
2002	9906	9759	9838	10665	11098	11919	12574	13132	13299	13647	13831	16389	17248	17665	18684	19373	20125	21218	22191	24187	25266	26172	28991	29232	29610	30264	31103	31557	32977	33353	34124	34418	34591	32966	36783
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2.1	63.1	61.1	60.1	59.1	58.1	57.1	55.1	52.1	50.1	48.1	47.1	46.1	45.1	44.1	43.1	42.1	39.1	37.1	36.1	35.1	34.1	33.1	30.1	29.1	27.1	26.1	25.1	24.1	20.1	19.1	18.1	16.1	15.1	13.1	11.2
SEQ ID No. 695	SEQ ID No. 3194	№	SEQ ID No. 3126	0 №	Б В	SEQ ID No. 3025	SEQ ID No. 2919	Ö. So.	Ю Мо	S Q	D No.	Θ No	Ö So	В	© No	Ю В	Ω	Ω	Б В	D No.	₽	Ö. So	Б В	D No	Ö N S	SEQ ID No. 1072	D No.	D No.	ID No.	Ю Мо	ID No.	D No.	Ю No.		D No.

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40073	43009	43425	44285	48882	49240	50938	52288	53340	53990	55366	26997	57608	58682	61016	61521	62223	62967	63917	62999	66918	68722	71343	71955	73159	74464	75396	76065	77588	78490	79331	80278	83893	85529	86043
38133	42674	42970	43401	44404	49022	49931	51047	52297	53466	54089	55687	57180	57786	58908	61261	61495	62317	63171	64253	66646	66923	68626	71452	72017	73382	74623	75598	76245	77714	78711	79487	80420	83988	85555
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Contig54 Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54
2622.1 2605.3	2620.1	2619.1	2616.1	139.5	2611.1	2609.2	2607.2	3489.2	1582.2	1581.4	1156.3	1153.1	1152.2	3488.1	3486.1	3485.1	3484.1	3483.3	1809.4	3481.2	3480.4	833.3	2116.2	499.2	500.2	501.2	502.3	503.3	3161.1	3160.1	1049.2	1050.4	2368.5	1778.5
SEQ ID No. 1084 SEQ ID No. 1076	SEQ ID No. 1083	SEQ ID No. 1081	SEQ ID No. 1080	SEQ ID No. 308	SEQ ID No. 1079	SEQ ID No. 1078	SEQ ID No. 1077	SEQ ID No. 1648	SEQ ID No. 432	SEQ ID No. 431	SEQ ID No. 160	SEQ ID No. 159	SEQ ID No. 158	SEQ ID No. 1647	SEQ ID No. 1646	SEQ ID No. 1645	SEQ ID No. 1644	SEQ ID No. 1643	SEQ ID No. 588	SEQ ID No. 1642	SEQ ID No. 1641	SEQ ID No. 3334	SEQ ID No. 757	SEQ ID No. 2659	SEQ ID No. 2667	SEQ ID No. 2672	SEQ ID No. 2677		SEQ ID No. 1435	SEQ ID No. 1434	SEQ ID No. 90	Ö. So	₽	SEQ ID No. 564

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39.2		٤	96578	90926	,	1	%86	•	
88.1		Ε	97597	98373	,	1	%66	ı	
3785.2		Ε	98361	98266	1	1	%66	ı	
34.2		Ε	99175	100311			%86		
33.1		Ε	100438	100734	,	1	94%	1	
2355.2	Contig54	E	100931	101824	1	ı	%26	1	
3780.2	Contig54	Ε	102158	102685	•	,	. 62%	•	
1505.4	Contig54	<u>a</u>	102899	105229		i	%96	. +	
1507.3	Contig54	a	105338	106840	ı	1	%86	1	
3454.1	Contig54	a.	106841	107404		ī	91%	+	
3453.1	Contig54	d	107695	109209		ı	%06	ı	
3451.1	Contig54	۵	109351	110169	,	1.	95%	+	
3450.2	Contig54	۵	110251	111336		ı	%86	•	
5135.2	Contig54	Ε	111413	112909		,	93%	+	
45.2	Contig54	Ε	113096	114103	ı	ı	%06	+	
2546.4	Contig54	٤	113946	115451		ı	94%		
3933.3	Contig54	d	115898	117001		1	%96	+	
1261.2	Contig54	Ω	117232	118920		ı	%66	•	
1264.3	Contig54	Ε	118978	120174	+	•	%56		
4485.2	Contig54	<u>α</u> .	120515	121345	+	•	%86	•	
4484.1	Contig54	۵	121446	122228	,	•	%26	+	
895.3	Contig54	ď	122576	123634		•	%86	•	
894.3	Contig54	٤	123736	124005		1	%86	+	
2220.2	Contig54	Q.	124301	124963		•	%66	•	
2219.2	Contig54	Ε	124648	125064		•	%86	•	
4319.2	Contig54	Ε	125030	125764	+	,	%86	+	

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	125926	127188	128626	120020	129491	129885	131646	132369	134827	136221	137374	197907	137907	139523	141103	142846	143553	144767	146263	147324	147869	149133	140702	143/02	151350	152533	157120	158050	159106	160503	161013	162074	102071	103332	164626	164875	165491	166111	
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166885	167598	168194	169230	170325	171626	172504	173129	174713	176087	176920	176950	177700	178381	178929	179417	180142	180883	181805	182843	183792	184053	45	205	983	1221	1823	2536	3956	4093	6821	7135	7683	8818	9401	10464
٤	E	Ε	Ε	Ε	Ε	Ε	a	a	E	٤	a	а	ď	٤	<u>α</u>	a	Ε	Ω.	<u>α</u>	۵	٤	۵	۵	۵	۵	۵	۵	۵	۵	d	a.	α.	d	ď	۵
Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig54	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55
853.1	852.3	3772.2	1101.4	1102.2	2255.2	3778.2	5550.3	5552.2	4409.1	4408.1	4407.1	4406.1	4405.1	4404.1	4403.2	4402.2	2598.3	5115.2	5116.3	5554.2	6018.1	5217.1	2659.2	2660.1	2661.2	2662.2	485.3	486.2	254.2	255.1	256.1	258.2	2006.2	1344.3	5219.1
ID No.	 	ID No.	SEQ ID No. 127	SEQ ID No. 128	SEQ ID No. 843		S O	SEQ ID No. 2948	⊡ So.	Ю В	ġ.	ID No.	ġ.	ġ	ġ	ġ	9	ું	9	ġ	ġ	ġ	SEQ ID No. 1107	9	ું	9	છું	ò	ું	SEQ ID No. 1038	SEQ ID No. 1045	SEQ ID No. 1058	D No.	_	SEQ ID No. 2786

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83683 84168 84934 85960 86924 87676 88268 89080 90691 91527 94135 96417 97093 97093 100433 101302 102432 102432 102432 105362 98262 99168 100904 102816 104112 105337 107019 107629 95677 96236 97094 85953 86954 87747 88340 93167

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	112373	113352	113977	115163	116406	117433	118891	119493	120958	122110	122374	123593	124882	125159	126202	127253	128618	128900	129671	130156	130624	131637	132518	133049	134262	136010	137527	138206	138772	139582	142158	142691	143953	145313	145939	147059
	E	Ξ	٤	۵	Ω	۵	۵	α.	Ε	a	. Δ.	۵	۵	۵	۵	۵	۵	. م	۵	۵	۵	α.	٤	٤	٤	d.	٤	ε	۵	Q.	۵	٤	Ε	Ε	ď	۵
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, , , ,	137.1	738.2	739.4	4947.2	4945.2	5078.2	5081.3	5080.4	5106.3	5108.2	5147.1	3500.3	3499.1	1226.2	1225.2	2275.3	2277.2	1171.3	1172.2	3498.1	616.4	591.3	5328.1	4209.2	2298.2	1035.3	4473.4	4474.1	4475.1	4477.2	4479.2	4480.4	5090.2	5088.2	5660.1	5056.2
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189518	191867	192574	193086	193707	194001	194995	196857	198794	199812	201254	202824	204695	205651	206564	207880	209053	210673	211888	212642	213318	215725	221632	220161	223283	224850	225843	226629	229195	230373	232594	233181	234512	235996	237023
188742	189807	191978	192553	193087	193708	194087	194986	196896	198790	199797	201664	202977	204692	205797	206522	207980	209183	210767	212031	212770	213377	215762	219910	221880	223444	224983	226063	226955	229465	230456	232591	233166	234482	236004
Ε	Ε	Ε	٤	Ε	Ε	٤	Ξ	Ε	ε	Ε	٤	٤	٤	۵	a	۵	۵	Ε	۵	۵	Ε	a	Ε	ε	E	Ε	۵	٤	۵	٥	٤	Ε	٤	Ε
Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contia55
1749.2	4179.2	1295.1	1294.2	1293.4	4337.3	4338.3	4339.1	4340.2	5386.1	4786.2	4785.1	4751.2	1516.3	1515.2	1513.4	5022.2	3988.3	3986.1	3985.1	856.2	857.2	4908.3	3996.2	1671.3	3995.5	3994.5	3993.2	887.4	5711.2	5193.2	3938.2	3937.2	2272.3	2274.2
SEQ ID No. 544	SEQ ID No. 2110	SEQ ID No. 252	SEQ ID No. 251	9	SEQ ID No. 2214	SEQ ID No. 2215	SEQ ID No. 2216	SEQ ID No. 2218	SEQ ID No. 2859	SEQ ID No. 2532	SEQ ID No. 2531	SEQ ID No. 2506	SEQ ID No. 387	SEQ ID No. 386	SEQ ID No. 385	SEQ ID No. 2679	SEQ ID No. 1967	SEQ ID No. 1966	SEQ ID No. 1965	SEQ ID No. 3348	SEQ ID No. 3349	SEQ ID No. 2612	SEQ ID No. 1974	SEQ ID No. 494					SEQ ID No. 3028	SEQ ID No. 2772	 	ID No.	Se Se	SEQ ID No. 855

SEQ ID No. 684	1972.3	Contig55	E	238224	239414	•	ı	%66	•	
SEQ ID No. 1186	278.3	Contig55	Ε	239267	241183		ı	%86	•	
D No.	5194.1	Contig55	Ε	241056	241625	,	+		•	
SEQ ID No. 1178	277.1	Contig55	Ε	241772	242491	ı	•	100%	'	
Ö. So	276.1	Contig55	Ε	242556	242792		ı	100%	,	
SEQ ID No. 547	1754.3	Contig55	Ε	243052	244590		1	%86	•	
D No	1755.3	Contig55	Ε	244529	244864		ı	%86	+	
SEQ ID No. 549	1756.4	Contig55	Ε	244943	245797	ı	ı	%86	1	
	2669.2	Contig55	Ε	246338	247576	ī	1	%66		
	2670.1	Contig55	٤	247577	248836		1	%66	1	
	2671.1	Contig55	đ	248990	249631	1	ı	%26	•	
	2674.1	Contig55	a	249701	252391			+ %66	ı	
ID No.	2677.2	Contig55	٩	252653	254050	+		100%	ı	
В В	847.4	Contig55	Ε	254244	257435	•	1	%66	ı	
SEQ ID No. 1121	2680.1	Contig55	۵	257935	258630	+	,	%66	1	
SEQ ID No. 518	1708.4	Contig55	۵	259050	259427			%66	`	
SEQ ID No. 519	1709.1	Contig55	Ε	259455	260177	,	ı	%86	•	
SEQ ID No. 520	1711.2	Contig55	Ε	260134	260889		ı	%66	1	
SEQ ID No. 864	2292.2	Contig55	۵	261111	262802		ı	%56	+	
SEQ ID No. 1122	2682.1	Contig55	a	262935	264578		ı	93%	+	
SEQ ID No. 80	1036.4	Contig55	Ε	264665	266002		ı	%86	•	
2 ID No. 81	1037.1	Contig55	Ε	265921	266478			100%	•	
SEQ ID No. 82	1039.2	Contig55	۵	266605	267834			95%	1	
	2144.3	Contig55	۵	267780	268817		•	91%	•	
ė	2683.3	Contig55	۵	268818	270377	,	•	%96	•	
Ю В	5195.2	Contig55	a	270293	271729		,	%86	•	
₽	2685.2	Contig55	۵	271681	272466			%86	+	
	4732.3	Contig55	Ε	272544	272921		•	%26	1	
ID No.	1743.4	Contig55	Ε	273331	273900		•	%66	•	
Ω So	1742.3	Contig55	Ε	273878	274645	,	,	%86	1	
S S	4733.2	Contig55	Ε	274618	275526		ı	%86	•	
D No.	4999.2	Contig55	Ε	275562	276383			%66	•	
D No.	5000.3	Contig55	۵	276648	277070	+	•	%86	•	
0 0 0	5001.1	Contig55	۵	277565	278038		ı	%66	+	
S Q	5307.3	Contig55	Ε	278103	279803	1	,	%26	1	
SEQ ID No. 1970	3991.3	Contig55	Ε	280182	281885			%36	•	

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282727	285299	285770	287344	288849	288577	289852	290118	292959	293819	294718	295850	296599	297459	298577	300040	302186	302744	304198	305635	306838	309283	309784	310592	311045	311026	311460	312241	312647	313828	314516	315411	316543	319145	317566	318523
281957	282705	285321	285761	287341	288191	288995	289828	290434	293013	293795	294663	295754	296578	297507	298751	300027	302427	302741	304199	305819	306980	309284	309795	310647	310700	311164	311846	312219	312599	313737	314611	315524	316557	316811	317987
Ω.	. a	Ε	Ε	Ε	a.	۵	۵	٤	E	E	Ε	Ε	٤	۵	٤	Ε	a	a	a	۵	۵	a	a	a.	a	Ε	Ε	٤	Ε	Ε	٥	E	Q.	ε	Ε
Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55	Contig55
837.2	835.2	3989.1	1210.2	1209.2	1208.1	5309.1	2514.3	893.4	4472.2	4744.3	1503.4	1501.3	2388.3	2387.3	1144.3	1145.3	5311.1	4094.2	4093.1	2343.2	2344.4	5478.2	1115.4	1113.2	4902.1	5477.1	5476.1	3556.2	3557.1	427.3	428.1	429.2	2092.2	2091.1	3558.1
SEQ ID No. 3336	SEQ ID No. 3335	D No.	SEQ ID No. 195	SEQ ID No. 193	SEQ ID No. 192	SEQ ID No. 2821		SEQ ID No. 3376	SEQ ID No. 2310	SEQ ID No. 2501	SEQ ID No. 378	SEQ ID No. 377	SEQ ID No. 928	SEQ ID No. 927	SEQ ID No. 154	SEQ ID No. 155	SEQ ID No. 2823	SEQ ID No. 2044	SEQ ID No. 2043	SEQ ID No. 893	SEQ ID No. 894	SEQ ID No. 2906	SEQ ID No. 136	SEQ ID No. 135	SEQ ID No. 2610	SEQ ID No. 2905	SEQ ID No. 2904	SEQ ID No. 1698	SEQ ID No. 1699	SEQ ID No. 2168	SEQ ID No. 2173	SEQ ID No. 2180	Ω So		SEQ ID No. 1700

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SEQ ID No. 1991	4016.1	Contig55	ε	357458	358642	1	,	%86	ı
SEQ ID No. 1992	4017.4	Contig55	٤	358638	360716		ı	%66	
SEQ ID No. 2848	5361.2	Contig55	Ε	360713	361897		•	%66	ı
SEQ ID No. 2847	5360.2	Contig55	۵	361044	361610		•	%66	+
	2592.4	Contig55	Ф	361844	362254		•	%26	+
₽	2593.3	Contig55	٤	361861	362634		1	%66	•
SEQ ID No. 2680	5026.1	Contig55	Ε	362635	362964	+	,	%86	•
SEQ ID No. 1068	2595.4	Contig55	E	362913	363662	+	,	%26	1
SEQ ID No. 2845	5359.2	Contig55	Ε	363663	364091		1	%86	ı
SEQ ID No. 2352	4526.3	Contig55	Ε	364070	364414	1	,	%96	
SEQ ID No. 434	1585.2	Contig55	Ε	364415	365494		,	%26	ı
SEQ ID No. 435	1586.2	Contig55	٤	365921	366190	+	,	100%	+
SEQ ID No. 436	1587.4	Contig55	a	366186	367574		ı	%26	ı
SEQ ID No. 2171	4275.2	Contig55	ď	367445	368293	+	1	%26	ı
SEQ ID No. 2170	4273.1	Contig55	d	368190	369320			%26	ı
SEQ ID No. 876	2313.2	Contig55	Ε	369399	370625		•	%96	ı
SEQ ID No. 875	2312.1	Contig55	٤	370829	371116		1	%86	+
SEQ ID No. 874	2311.4	Contig55	٤	371117	373702		1	%86	
SEQ ID No. 627	1873.2	Contig55	Ε	373717	374169		1	100%	•
SEQ ID No. 626	1871.2	Contig55	Ε	374162	375211		1	100%	•
SEQ ID No. 2428	4643.2	Contig55	Ε	375511	376461		1	%66	+
SEQ ID No. 2427	4642.1	Contig55	Ε	376740	377243		•	+ %86	
SEQ ID No. 3101	5910.1	Contig55	٤	377311	377523			%86	+
SEQ ID No. 2426	4640.2	Contig55	a	377831	380395	1	1	%66	•
SEQ ID No. 1994	4020.2	Contig55	م	380396	382096		ı	%66	
SEQ ID No. 3332	830.4	Contig55	۵	382039	384633			%36	+
SEQ ID No. 3333	832.3	Contig55	۵	384667	385662	ı	ı	94%	1
SEQ ID No. 3041	5747.1	Contig55	۵	385787	386191	+	•	%96	+
SEQ ID No. 678	1955.3	Contig55	Ε	386532	388040	+		95%	
SEQ ID No. 885	2328.2	Contig55	۵	388278	390869	•	1	%66	+
SEQ ID No. 1501	3269.1	Contig55	۵	390859	394104		ı	%86	ı
SEO ID No. 960	2430.2	Contig55	۵	394309	395385		•	%66	1
SEQ ID No. 1500	3267.1	Contig55	Ω.	395435	396055		1	%66	+
Ω	3265.1	Contig55	Ε	396103	396999	,		%86	Ī
Ω So	3263.2	Contig55	۵	397202	398425	•	•	%86	•
SEQ ID No. 1497	3260.2	Contig55	Ε	398427	399434	+	ı	%66	+

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SEO ID No. 3040	5743.1	Contin55	٤	435125	435313		+			٠ +
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ID No.	3744.2	Contig56	۵	629	1834		+			+
SEQ ID No. 1828	3743.1	Contig56	Ε	1999	2334	+	Ī	%96		+
SEQ ID No. 1827	3742.1	Contig56	Ε	2352	2624		1	95%		+
	3741.1	Contig56	Ε	2635	3855	1	•	%86		+
	3740.3	Contig56	Ε	4089	4625	,	•	100%		
So O	3739.3	Contig56	۵	4848	5291	1	ı	%66		
	3737.1	Contig56	a.	5344	6447	+	1	94%		+
D No.	1082.3	Contig56	٤	6468	8033	1	+			
SEQ ID No. 357	1469.4	Contig56	٤	8879	12292	ı	+			+
D No.	4312.2	Contig56	Ε	12373	13725	1	•	%86		
SEQ ID No. 2196	4311.1	Contig56	٥	13958	14506		1	%96		
о В	4310.1	Contig56	Ε	14610	15161		,	%26		+
D No.	4309.2	Contig56	٤	15077	16219		•	%26		<u>,</u> +
SEQ ID No. 2010	4047.3	Contig56	٤	16291	17385	+	ı	%66		•
SEQ ID No. 2011	4048.3	Contig56	۵	17493	19334		ı	100%		ı
SEQ ID No. 2013	4050.1	Contig56	۵	19403	20197	+	1	%86		1
SEQ ID No. 2014	4051.2	Contig56	۵	20206	20601	+	1	%86		ı
SEQ ID No. 2015	4054.2	Contig56	۵	20585	21265			%66		
SEQ ID No. 2016	4055.1	Contig56	٤	21293	23185		+		+	
SEQ ID No. 2017	4056.2	Contig56	Ε	23266	24495		ı	%66		1
SEQ ID No. 2018	4058.2	Contig56	Ε	24583	24942		+			+
SEQ ID No. 472	1641.5	Contig56	۵	25345	26694		1	%86		
SEQ ID No. 473	1642.3	Contig56	a	26679	27341		1	100%		
SEQ ID No. 2465	4699.3	Contig56	a.	27452	28756		,	%66		
SEQ ID No. 2340	4510.1	Contig56	<u>a</u>	28978	31440			100%		ı
SEQ ID No. 2341	4511.1	Contig56	۵	31527	31847		•	%86	+	ı
ġ	4512.4	Contig56	a	32084	33979		ı	%86		1
9	1787.4	Contig56	۵	34519	37686		+			+
	3530.1	Contig56	Ε	37969	38598		+			+
SEQ ID No. 1679	3532.1	Contig56	a	38652	41942		+			+
Ω	1487.3	Contig56	Ε	42394	43230			%66	+	
Ω	3533.1	Contig56	Ε	43296	44021	+	•	%66		1
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SEQ ID No. 2797	9.	SEO ID No. 343	SEQ ID No. 625	SEQ ID No. 3195	SEQ ID No. 1460	SEQ ID No. 1461	₽	\Box		SEO ID No 190	SEO ID No 191	SEQ ID No. 1280	SEO ID No 1282	SEO ID No. 1283	SEQ ID No. 1284	SEO ID No. 813	SEO ID No. 944	SEC ID NO. 814	SECULD INO. 815	<u>.</u>	SEC ID No. 1043	SEQ ID No. 1042	SEQ ID No. 2408	SEQ ID No. 2407	9 1 1	SEQ ID No. 2405	Q	ю В	ġ	D So.	₽	Ω	Θ S O	<u>0</u> №	SEC ID No 2007

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432873	435357	436686	437445	437537	440810	442423	443439	443846	444206	446531	447359	448649	450084	451509	452294	453233	453914	455395	456141	456543	456954	458774	460258	461427	462305	462634	463276	463899	465296	465798	468423	469441	470268	470001
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2268.2	2264.2	1242.2	1239.2	1238.2	5188.2	5189.2	3235.3	5947.1	3236.3	926.4	928.1	929.2	1788.3	1792.2	3238.1	1116.2	1118.2	1119.2	3239.1	3240.3	4192.2	4191.1	4188.1	4186.1	4184.1	1647.2	1645.1	1644.6	2583.3	2582.3	4809.2	4810.1	4811.2	1768.3
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Contig56 4071.1 4070.3 4678.3 4680.2 4683.3 5492.2 4356.2 4356.2 4354.2 4354.2 4354.2 4354.2 4354.2 4354.2 189.4 1331.3 1331.3 1331.3 1322.3 3625.2 3622.3 4977.3 4977.3 4977.3 4977.3 SEQ ID No. 2562
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510713	512598	513518	514374	515128	515479	518054	519483	520040	520200	521370	527089	528497	529698	530771	531595	533008	534412	536875	537264	537425
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327.2	328.1	330.2	2920.1	2919.1	1003.3	1005.2	2917.1	2916.1	2360.2	2361.4	3568.3	3566.1	712.2	711.2	709.1	2335.2	2336.2	3564.2	3563.2	5988.1
SEQ ID No. 1502	SEQ ID No. 1510	SEQ ID No. 1523	SEQ ID No. 1264	SEQ ID No. 1262	SEQ ID No. 61	SEQ ID No. 62	SEQ ID No. 1261	SEQ ID No. 1260	SEQ ID No. 909	SEQ ID No. 910	SEQ ID No. 1707	SEQ ID No. 1706	SEQ ID No. 3251	SEQ ID No. 3250	SEQ ID No. 3249	SEQ ID No. 888	SEQ ID No. 889	SEQ ID No. 1705	SEQ ID No. 1704	SEQ ID No. 3123

ORF Best-BlastP

- Best-BlastP=> >nrprot 47% Identities = 35/90 (38%), Positives = 55/90 (61%), Gaps = 5/90 (5%) dbj|BAC94688.1| hypothetical protein [Vibrio Length = 343 /ulnificus YJ016] 10.1
- Best-BlastP=> >nrprot 62% Identities = 120/257 (46%), Positives = 162/257 (63%), Gaps = 3/257 (1%) ref|ZP_00079402.1| COG1024: Enoyl-1000.2
 - Best-BlastP=> >nrprot 97% Identities = 419/437 (95%), Positives = 425/437 (97%) gb|AAM00614.1| chemiosmotic efflux system protein C-like Length = 272metallireducens CoA hydratase/carnithine racemase [Geobacter 1001.3
 - Best-BlastP=> >nrprot 11% Identities = 35/129 (27%), Positives = 64/129 (49%), Gaps = 6/129 (4%) ref[ZP_00108772.1| COG0642: Signal Length = 2053 Length = 448 transduction histidine kinase [Nostoc punctiforme] pneumophila] 1002.2
 - 1003.3 Best-BlastP=> >nrprot No Hits found
- Best-BlastP=> >nrprot 67% Identities = 126/243 (51%), Positives = 165/243 (67%), Gaps = 4/243 (1%) ref[NP_820690.1] dihydrodipicolinate reductase [Coxiella burnetii RSA 493] sp|P24703|DAPB_COXBU Dihydrodipicolinate reductase (DHPR) gb|AA091204.1| dihydrodipicolinate Length = 239 reductase [Coxiella burnetii RSA 493] 1005.2
- OXIDOREDUCTASE PROTEIN [Ralstonia solanacearum] emb|CAD15760.1| OXIDOREDUCTASE PROTEIN [Ralstonia solanacearum] Best-BlastP=> >nrprot 61% Identities = 96/211 (45%), Positives = 135/211 (63%), Gaps = 18/211 (8%) ref[NP_520174.1| PROBABLE PROBABLE TRANSMEMBRANE NADH DEHYDROGENASE I (CHAIN J) TRANSMEMBRANE NADH DEHYDROGENASE I (CHAIN J) Length = 210 1006.3
- 9a5c] ref[ZP_00039600.1| hypothetical protein [Xylella fastidiosa Dixon] ref[ZP_00041894.1| hypothetical protein Xylella fastidiosa (strain 9a5c) gb|AAF83124.1|AE003884_9 NADH-Best-BlastP=> >nrprot 79% Identities = 113/159 (71%), Positives = 132/159 (83%) refINP_297604.1| NADH-ubiquinone oxidoreductase, NQO9 Temecula1] pir||C82822 9a5c] gb|AAO28142.1| NADH-ubiquinone oxidoreductase NQO9 subunit Xylella fastidiosa Ann-1] ref|NP_778493.1| NADH-ubiquinone oxidoreductase NQO9 subunit [Xylella fastidiosa NADH-ubiquinone oxidoreductase, NQO9 subunit XF0313 [imported] ubiquinone oxidoreductase, NQO9 subunit [Xylella fastidiosa Length = 163 Temecula1] subunit [Xylella fastidiosa 1009.2
- Best-BlastP=> >nrprot 82% Identities = 223/332 (67%), Positives = 283/332 (85%) refINP_820424.1| NADH dehydrogenase I, H subunit [Coxiella Length = 340ournetii RSA 493] gb|AAO90938.1| NADH dehydrogenase I, H subunit [Coxiella burnetii RSA 493] 1010.2
 - dehydrogenase I, G subunit [Coxiella burnetii RSA 493] gb|AAO90939.1| NADH dehydrogenase I, G subunit [Coxiella burnetii RSA 493] Identities = 374/800 (46%), Positives = 511/800 (63%), Gaps = 32/800 (4%) ref[NP_820425.1| NADH Best-BlastP=> >nrprot 65% 1011.5
- 1013.1 Best-BlastP=> >nrprot No Hits found
- 3est-BlastP=> >nrprot 55% Identities = 163/485 (33%), Positives = 254/485 (52%), Gaps = 36/485 (7%) ref[NP_711514.1] putative amidase Leptospira interrogans serovar lai str. 56601] gb/AAN48532.1/AE011313_8 putative amidase [Leptospira interrogans serovar lai str. 56601] Length = 5001014.2
- Identities = 811/842 (96%), Positives = 830/842 (98%) gb|AAM00631.1| putative cation efflux transporter [Legionella Length = 842 3est-BlastP=> >nrprot 92% 102.1

- Identities = 588/1008 (58%), Positives = 775/1008 (76%), Gaps = 4/1008 (0%) ref[NP 820092.1] transporter, AcrB/AcrD/AcrF family [Coxiella burnetii RSA 493] gb|AAO90606.1| transporter, AcrB/AcrD/AcrF family [Coxiella burnetii RSA 493] Best-BlastP=> >nrprot 76% 1021.3
- A2012] ref[NP_844481.1| acetyltransferase, GNAT family [Bacillus anthracis str. Ames] Identities = 76/278 (27%), Positives = 135/278 (48%), Gaps = 6/278 (2%) ref|NP_655937.1| Acetyltransf, Length = 295gb|AAP25967.1| acetyltransferase, GNAT family [Bacillus anthracis str. Ames] Acetyltransferase (GNAT) family [Bacillus anthracis Best-BlastP=> >nrprot 47% 1022.2
- Best-BlastP=> >nrprot 65% Identities = 89/178 (50%), Positives = 126/178 (70%), Gaps = 3/178 (1%) ref[NP_799717.1] peptide methionine RIMD 2210633] dbj|BAC61550.1| peptide methionine sulfoxide reductase [Vibrio sulfoxide reductase [Vibrio parahaemolyticus Length = 380parahaemolyticus] 1023.2
- Best-BlastP=> >nrprot 54% Identities = 71/193 (36%), Positives = 108/193 (55%), Gaps = 2/193 (1%) ref[NP_561497.1| conserved hypothetical Length = 195 protein [Clostridium perfringens] dbj|BAB80287.1| conserved hypothetical protein [Clostridium perfringens str. 13]

1024.2

- Identities = 221/382 (57%), Positives = 274/382 (71%), Gaps = 2/382 (0%) ref[ZP_00032722.1| COG1228: Length = 431 [Burkholderia fungorum] midazolonepropionase and related amidohydrolases Best-BlastP=> >nrprot 67% 1028.3
- Identities = 414/419 (98%), Positives = 418/419 (99%) gb/AAM00630.1| chemiosmotic efflux system B protein C Length = 419 Best-BlastP=> >nrprot 99% [Legionella pneumophila] 103.1

1030.3

- Identities = 70/234 (29%), Positives = 115/234 (49%), Gaps = 6/234 (2%) refINP_867781.1| probable sepiapterin Length = 261reductase homolog yueD [Pirellula sp.] emb|CAD75328.1| probable sepiapterin reductase homolog yueD [Pirellula sp.] Best-BlastP=> >nrprot 45% 1032.3
- Identities = 131/317 (41%), Positives = 188/317 (59%), Gaps = 16/317 (5%) refINP_520766.1| PROBABLE ARGINASE PROTEIN [Ralstonia solanacearum] emb|CAD16352.1| PROBABLE ARGINASE PROTEIN [Ralstonia solanacearum] Best-BlastP=> >nrprot 58%
- Identities = 43/131 (32%), Positives = 69/131 (52%), Gaps = 8/131 (6%) gb|AAL78307.1|AF288617_4 Dotl Length = 212 Best-BlastP=> >nrprot 21% Legionella longbeachae] 1034.3
- Identities = 124/492 (25%), Positives = 207/492 (42%), Gaps = 95/492 (19%) ref[NP_487180.1] hypothetical protein 7120) dbj|BAB74839.1| Nostoc sp. PCC 7120] pir[|AE2198 hypothetical protein alr3140 [imported] - Nostoc sp. (strain PCC Length = 455 ORF_ID:alr3140~hypothetical protein [Nostoc sp. PCC 7120] Best-BlastP=> >nrprot 43% 1035.3
- Identities = 206/421 (48%), Positives = 258/421 (61%), Gaps = 8/421 (1%) ref[ZP_00043700.1| COG0389: Length = 421 repair [Magnetococcus sp. MC-1] Nucleotidyltransferase/DNA polymerase involved in DNA Best-BlastP=> >nrprot 61% 1036.4
- Identities = 71/123 (57%), Positives = 96/123 (78%) ref|ZP_00043701.1| COG1974: SOS-response transcriptional Length = 238 autopeptidases) [Magnetococcus sp. MC-1] Best-BlastP=> >nrprot 68% epressors (RecA-mediated 1037.1
 - Identities = 177/396 (44%), Positives = 241/396 (60%), Gaps = 5/396 (1%) ref|ZP_00094397.1| COG0624: desuccinylase and related deacylases [Novosphingobium Acetylornithine deacetylase/Succinyl-diaminopimelate Best-BlastP=> >nrprot 59% -ength = 465 1039.2
- Identities = 371/377 (98%), Positives = 376/377 (99%) gb|AAM00629.1| chemiosmotic efflux system B protein B Length = 377 Best-BlastP=> >nrprot 99% Legionella pneumophila] 104.1

str. DC3000] gb[AAO58815.1] conserved hypothetical protein [Pseudomonas Best-BlastP=> >nrprot 63% Identities = 150/366 (40%), Positives = 220/366 (60%), Gaps = 25/366 (6%) ref[NP_795120.1| conserved Length = 368 hypothetical protein [Pseudomonas syringae pv. tomato str. DC3000] syringae pv. tomato 1040.3

1041.3

Identities = 557/801 (69%), Positives = 663/801 (82%), Gaps = 1/801 (0%) ref[NP_756480.1| DNA gyrase subunit B Length = 805 Escherichia coli CFT073] gb|AAN83054.1|AE016769_169 DNA gyrase subunit B [Escherichia coli CFT073] Best-BlastP=> >nrprot 82%

1042.5

Best-BlastP=> >nrprot 76% Identities = 264/406 (65%), Positives = 314/406 (77%) ref[NP_719647.1] malate oxidoreductase, putative Shewanella oneidensis MR-1] gb|AAN57091.1|AE015843_3 malate oxidoreductase, putative [Shewanella oneidensis MR-1]

1044.3

Identities = 152/394 (38%), Positives = 236/394 (59%), Gaps = 12/394 (3%) ref[NP_820159.1| major facilitator family ransporter [Coxiella burnetii RSA 493] gb|AAO90673.1| major facilitator family transporter [Coxiella burnetii RSA 493] 3est-BlastP=> >nrprot 59%

3est-BlastP=> >nrprot 59% Identities = 142/261 (54%), Positives = 179/261 (68%), Gaps = 1/261 (0%) ref[ZP_00102270.1| COG1075: Predicted Length = 364 alpha/beta hydrolase fold [Desulfitobacterium hafniense] acetyltransferases and hydrolases with the

1047.2

Identities = 33/98 (33%), Positives = 48/98 (48%), Gaps = 3/98 (3%) ref[NP_717311.1| hypothetical protein Lenath = 443 Shewanella oneidensis MR-1] gb[AAN54755.1|AE015616_1 hypothetical protein [Shewanella oneidensis MR-1] Best-BlastP=> >nrprot 12% 1048.3

Best-BlastP=> >nrprot 48% Identities = 74/232 (31%), Positives = 112/232 (48%), Gaps = 10/232 (4%) refINP_069069.1| glutamine ABC

1049.2

(glnH) homolog - Archaeoglobus fulgidus gb[AAB91001.1] glutamine ABC transporter, periplasmic (glnH) [Archaeoglobus fulgidus DSM 4304] pirl|G69278 glutamine ABC transporter, Length = 264(glnH) [Archaeoglobus fulgidus DSM 4304] ransporter, periplasmic glutamine-binding protein periplasmic glutamine-binding protein glutamine-binding protein

Identities = 96/410 (23%), Positives = 164/410 (40%), Gaps = 57/410 (13%) pir||T31688 Ca2+-transporting ATPase Identities = 86/174 (49%), Positives = 114/174 (65%), Gaps = 2/174 (1%) ref[NP_820971.1] phosphatase, putative tetraurelia gb|AAB81284.1| plasma membrane calcium ATPase [Paramecium tetraurelia] EC 3.6.3.8), plasma membrane - Paramecium Best-BlastP=> >nrprot 64% Best-BlastP=> >nrprot 14% 1050.4

Coxiella burnetii RSA 493] gb|AAO91485.1| phosphatase, putative [Coxiella burnetii RSA 493] 1053.2

Best-BlastP=> >nrprot 59% Identities = 102/256 (39%), Positives = 144/256 (56%), Gaps = 16/256 (6%) ref|NP_820853.1| conserved lypothetical protein [Coxiella burnetii RSA 493] gb[AAO91367.1] conserved hypothetical protein [Coxiella burnetii RSA 493] 1055.1

Length = 184

Best-BlastP=> >nrprot No Hits found 1056.1

discoideum) gb|AAA33227.1| myosin Best-BlastP=> >nrprot 43% Identities = 112/511 (21%), Positives = 204/511 (39%), Gaps = 94/511 (18%) sp|P08799|MYS2_DICDI Myosin II neavy chain, non muscle pir||A26655 myosin heavy chain [similarity] - slime mold (Dictyostelium 1058.3

Identities = 302/423 (71%), Positives = 364/423 (86%), Gaps = 2/423 (0%) ref|NP_820394.1| citrate synthase Coxiella burnetii RSA 493] sp|P18789|CISY_COXBU Citrate synthase pir||JQ1392 citrate (si)-synthase (EC 4.1.3.7) - Coxiella burnetii Length = 430 gb|AAA23307.1| citrate synthase (g1tA) (EC 4.1.3.7) gb|AAO90908.1| citrate synthase [Coxiella burnetii RSA 493] 3est-BlastP=> >nrprot 85%

Identities = 114/286 (39%), Positives = 168/286 (58%), Gaps = 7/286 (2%) ref[NP_813465.1] purine nucleoside VPI-5482] gb|AAO79659.1| purine nucleoside phosphorylase II [Bacteroides Length = 292 phosphorylase II [Bacteroides thetaiotaomicron VPI-5482] Best-BlastP=> >nrprot 57% thetaiotaomicron 1060.2

1063.1 Best-BlastP=> >nrprot No Hits found

Best-BlastP=> >nrprot 48% Identities = 124/365 (33%), Positives = 187/365 (51%), Gaps = 10/365 (2%) dbj|BAA31547.1| metal-activated Length = 379pyridoxal enzyme [Arthrobacter sp.] 1065.3

1066.2

Best-BlastP=> >nrprot 53% Identities = 124/294 (42%), Positives = 160/294 (54%), Gaps = 18/294 (6%) ref|ZP_00016110.1| COG0596: Length = 286 hydrolase superfamily) [Rhodospirillum rubrum] Predicted hydrolases or acyltransferases (alpha/beta

Identities = 28/70 (40%), Positives = 40/70 (57%), Gaps = 11/70 (15%) ref[NP_489272.1] unknown protein [Nostoc 7120) dbj|BAB76931.1| sp. PCC 7120] pir[|AH2459 hypothetical protein alr5232 [imported] - Nostoc sp. (strain PCC Length = 204ORF_ID:alr5232~unknown protein [Nostoc sp. PCC 7120] Best-BlastP=> >nrprot 47% 1067.1

Best-BlastP=> >nrprot 46% Identities = 103/273 (37%), Positives = 145/273 (53%), Gaps = 35/273 (12%) emb|CAA75849.1| hypothetical protein Length = 309Coxiella burnetii] 1069.2

Identities = 1034/1047 (98%), Positives = 1041/1047 (99%) gb|AAM00628.1| chemiosmotic efflux system B protein A Length = 1047 Best-BlastP=> >nrprot 99% Legionella pneumophila] 107.1

Identities = 411/418 (98%), Positives = 416/418 (99%) pir||T08882 proline/betaine transport protein homolog CitA pneumophila emb|CAA75171.1| TphA protein [Legionella pneumophila] gb|AAC38182.1| CitA [Legionella pneumophila] Length = 418 emb|CAA75337.1| TphA protein [Legionella pneumophila] Best-BlastP=> >nrprot 99% 1071.3

Identities = 966/973 (99%), Positives = 971/973 (99%) pirl|T18341 icmF protein - Legionella pneumophila emb|CAA75172.1| IcmF protein [Legionella pneumophila] emb|CAA75338.1| IcmF protein [Legionella pneumophila] Best-BlastP=> >nrprot 99% 1072.4

Identities = 343/638 (53%), Positives = 456/638 (71%), Gaps = 6/638 (0%) ref[NP_819606.1| fatty oxidation complex, alpha subunit [Coxiella burnetii RSA 493] gb|AAO90120.1 | fatty oxidation complex, alpha subunit [Coxiella burnetii RSA 493] Best-BlastP=> >nrprot 67% Length = 642 1074.3

Identities = 271/422 (64%), Positives = 333/422 (78%), Gaps = 1/422 (0%) emb|CAD58320.1| beta-Subunit of fatty Length = 426 3-keto-acyl-CoA-thiolase [Azoarcus sp. EbN1] Best-BlastP=> >nrprot 75% acid oxidation complex, 1075.3

12472] gb[AAQ60013.1] conserved hypothetical protein [Chromobacterium violaceum ATCC Identities = 50/119 (42%), Positives = 70/119 (58%) ref[NP_902011.1] conserved hypothetical protein Chromobacterium violaceum ATCC Best-BlastP=> >nrprot 56% Length = 122 1076.3

TRANSCRIPTION REGULATOR PROTEIN [Sinorhizobium meliloti] emb|CAC41614.1| PUTATIVE TRANSCRIPTION REGULATOR PROTEIN Identities = 137/226 (60%), Positives = 175/226 (77%), Gaps = 1/226 (0%) ref|NP_384333.1| PUTATIVE Length = 230 Best-BlastP=> >nrprot 75% Sinorhizobium meliloti] 1077.1

- Best-BlastP=> >nrprot 33% Identities = 65/262 (24%), Positives = 110/262 (41%), Gaps = 21/262 (8%) refINP_624872.1| hypothetical protein SCF73.06c [Streptomyces coelicolor A3(2)] emb[CAB57411.1| hypothetical protein SCF73.06c [Streptomyces coelicolor A3(2)] 1078.2
- Identities = 71/229 (31%), Positives = 113/229 (49%), Gaps = 11/229 (4%) ref[ZP_00081514.1| COG1560: Length = 324metallireducens] .auroyl/myristoyl acyltransferase [Geobacter Best-BlastP=> >nrprot 40% 1080.2
- Best-BlastP=> >nrprot 36% Identities = 81/362 (22%), Positives = 140/362 (38%), Gaps = 52/362 (14%) ref|ZP_00133556.1| COG5295: 1081.2
- Identities = 236/481 (49%), Positives = 325/481 (67%), Gaps = 12/481 (2%) ref[NP_484210.1] alpha,alpha-trehalase Nostoc sp. PCC 7120] pir||AF1827 alpha,alpha-trehalase [imported] - Nostoc sp. (strain PCC 7120) dbj|BAB77690.1| alpha,alpha-trehalase Length = 3391Autotransporter adhesin [Haemophilus somnus 2336] Length = 495 Best-BlastP=> >nrprot 67% Nostoc sp. PCC 7120] 1082.3
- Identities = 191/581 (32%), Positives = 316/581 (54%), Gaps = 34/581 (5%) ref[NP_488098.1| similar to isovaleryl-CoA dehydrogenase [Nostoc sp. PCC 7120] pir||AC2313 hypothetical protein alr4058 [imported] - Nostoc sp. (strain PCC Best-BlastP=> >nrprot 56% 1087.2
 - Best-BlastP=> >nrprot 40% Identities = 124/450 (27%), Positives = 202/450 (44%), Gaps = 37/450 (8%) ref|NP_870622.1| probable auxin-Length = 602 PCC 7120] dbj|BAB75757.1| ORF_ID:alr4058~similar to isovaleryl-CoA dehydrogenase [Nostoc sp. 1088.3
- Best-BlastP=> >nrprot 48% Identities = 274/902 (30%), Positives = 444/902 (49%), Gaps = 54/902 (5%) ref|ZP_00123122.1| hypothetical protein Length = 559 esponsive-like protein [Pirellula sp.] emb|CAD77699.1| probable auxin-responsive-like protein [Pirellula sp.] Length = 948 Haemophilus somnus 129PT] 1089.2
- Best-BlastP=> >nrprot 96% Identities = 441/469 (94%), Positives = 456/469 (97%) gb|AAM00627.1| unknown [Legionella pneumophila] Length = 470 109.1
- Identities = 28/84 (33%), Positives = 51/84 (60%), Gaps = 12/84 (14%) ref|ZP_00025260.1| hypothetical protein Length = 241 Best-BlastP=> >nrprot 43% Ralstonia metallidurans] 1090.1
 - Identities = 114/438 (26%), Positives = 192/438 (43%), Gaps = 79/438 (18%) refINP_873428.1| conserved nypothetical protein [Haemophilus ducreyi 35000HP] gb|AAP95817.1| conserved hypothetical protein [Haemophilus ducreyi 35000HP] Best-BlastP=> >nrprot 45% _ength = 481 1091.2
- Best-BlastP=> >nrprot 46% Identities = 64/221 (28%), Positives = 115/221 (52%), Gaps = 8/221 (3%) ref|ZP_00087883.1| COG1940: Length = 27fluorescens PfO-1] ranscriptional regulator/sugar kinase [Pseudomonas 1092.3
- Identities = 161/325 (49%), Positives = 232/325 (71%), Gaps = 16/325 (4%) refINP_819394.1| sohB protein [Coxiella Length = 338ournetii RSA 493] gb/AAO89908.1| sohB protein [Coxiella burnetii RSA 493] Best-BlastP=> >nrprot 74% 1093.3
 - Identities = 30/50 (60%), Positives = 34/50 (68%), Gaps = 4/50 (8%) gb|EAA17424.1| putative HSP protein Length = 929 Best-BlastP=> >nrprot 36% Plasmodium yoelii yoelii] 1094.1
- Identities = 253/483 (52%), Positives = 333/483 (68%), Gaps = 17/483 (3%) ref[ZP_00127277.1| COG2204; ATPase, and DNA-binding domains [Pseudomonas syringae Response regulator containing CheY-like receiver, AAA-type 3est-BlastP=> >nrprot 70% 1096.3

| Identities = 136/330 (41%), Positives = 195/330 (59%), Gaps = 3/330 (0%) refINP_718989.1 | conserved hypothetical protein [Shewanella oneidensis MR-1] sp|Q8EBR4|TRUD_SHEON tRNA pseudouridine synthase D (Pseudouridylate synthase) (Uracil gb|AAN56433.1|AE015780_4 conserved hypothetical protein [Shewanella oneidensis MR-1] Best-BlastP=> >nrprot 57%

1098.3

Best-BlastP=> >nrprot 63% Identities = 158/328 (48%), Positives = 207/328 (63%), Gaps = 6/328 (1%) ref|NP_403834.1| conserved hypothetical protein [Yersinia pestis] ref|NP_671254.1| hypothetical protein [Yersinia pestis KIM] pir||AH0022 conserved hypothetical protein YPO0180 (strain CO92) emb|CAC89042.1| conserved hypothetical protein [Yersinia pestis CO92] gb|AAM87505.1|AE014000_9 hypothetical protein [Yersinia pestis KIM] imported] - Yersinia pestis 1099.2

Identities = 77/276 (27%), Positives = 127/276 (46%), Gaps = 29/276 (10%) ref|ZP_00023112.1| hypothetical protein Length = 326 Best-BlastP=> >nrprot 39% Ralstonia metallidurans] 11.2

Identities = 128/359 (35%), Positives = 200/359 (55%), Gaps = 24/359 (6%) ref|NP_878747.1| histidinol-phosphate floridanus] emb|CAD83523.1| histidinol-phosphate aminotransferase [Candidatus Length = 356Length = 348 aminotransferase [Candidatus Blochmannia floridanus] Best-BlastP=> >nrprot 54% Blochmannia 1101.4

Best-BlastP=> >nrprot 66% Identities = 203/427 (47%), Positives = 287/427 (67%), Gaps = 9/427 (2%) dbj|BAC94115.1| histidinol Length = 431 dehydrogenase [Vibrio vulnificus YJ016] 1102.2

1105.2

Best-BlastP=> >nrprot 36% Identities = 187/439 (42%), Positives = 279/439 (63%), Gaps = 1/439 (0%) ref|NP_715981.1| sensory box protein [Shewanella oneidensis MR-1] gb[AAN53426.1|AE015481_9 sensory box protein [Shewanella oneidensis MR-1]

Identities = 1366/1368 (99%), Positives = 1367/1368 (99%) gb|AAC69338.1| RNA polymerase B-subunit [Legionella Length = 1368 Best-BlastP=> >nrprot 99% 1106.5

[Bacillus anthracis A2012] ref[NP_845551.1| 3-oxoacyl-(acyl-carrier-protein) synthase Best-BlastP=> >nrprot 66% Identities = 154/324 (47%), Positives = 219/324 (67%), Gaps = 3/324 (0%) ref|NP_657119.1| Chal_stil_syntC, anthracis str. Ames] gb/AAP27037.1| 3-oxoacyl-(acyl-carrier-protein) synthase III, putative [Bacillus Chalcone and stilbene synthases, C-terminal domain Length = 329III, putative [Bacillus 1107.2

[Bacillus anthracis str. Ames] gb[AAP27035.1] 3-beta hydroxysteroid dehydrogenase/isomerase Best-BlastP=> >nrprot 64% Identities = 150/329 (45%), Positives = 212/329 (64%), Gaps = 3/329 (0%) ref|NP_657117.1| 3Beta_HSD, 3-beta [Bacillus anthracis A2012] ref[NP_845549.1| 3-beta hydroxysteroid Length = 328 [Bacillus anthracis str. Ames] hydroxysteroid dehydrogenase/isomerase family dehydrogenase/isomerase family protein 1109.3

Best-BlastP=> >nrprot 97% Identities = 485/504 (96%), Positives = 493/504 (97%) splQ8RNP4|TYPH_LEGPN Putative thymidine phosphorylase Length = 517 (TdRPase) gb|AAM00626.1| unknown [Legionella pneumophila] 111.2

Best-BlastP=> >nrprot 58% Identities = 109/270 (40%), Positives = 164/270 (60%), Gaps = 6/270 (2%) ref[NP_657116.1] hypothetical protein A2012] ref[NP_845548.1| conserved hypothetical protein [Bacillus anthracis str. Ames] Length = 283gb|AAP27034.1| conserved hypothetical protein [Bacillus anthracis str. Ames] predicted by GeneMark [Bacillus anthracis 1111.3

Length = 116 Best-BlastP=> >nrprot 31% Identities = 37/58 (63%), Positives = 42/58 (72%), Gaps = 3/58 (5%) ref|NP_759576.1| Conserved hypothetical protein [Vibrio vulnificus CMCP6] gbJAAO09103.1|AE016799_1 Conserved hypothetical protein [Vibrio vulnificus CMCP6]

- str. 56601] gb|AAN47643.1|AE011231_6 conserved hypothetical protein [Leptospira interrogans Best-BlastP=> >nrprot 26% Identities = 41/132 (31%), Positives = 70/132 (53%), Gaps = 10/132 (7%) ref[NP_710625.1] conserved hypothetical Length = 359protein [Leptospira interrogans serovar lai str. 566011 1115.4
- 1116.2 Best-BlastP=> >nrprot No Hits found
- Identities = 277/452 (61%), Positives = 359/452 (79%), Gaps = 1/452 (0%) gb|AAM00627.1| unknown [Legionella Length = 470 Best-BlastP=> >nrprot 79% pneumophilal 1118.2
- 1119.2 Best-BlastP=> >nrprot No Hits found
- Identities = 142/291 (48%), Positives = 196/291 (67%), Gaps = 5/291 (1%) ref[NP_716239.1] hflC protein Length = 297 Shewanella oneidensis MR-1] gb/AAN53684.1/AE015507_10 hflC protein [Shewanella oneidensis MR-1] Best-BlastP=> >nrprot 64% 1122.1
- Identities = 176/379 (46%), Positives = 237/379 (62%), Gaps = 24/379 (6%) ref[NP_253629.1| protease subunit HflK (strain PAO1) Pseudomonas aeruginosa PA01] pir||B83028 proteinase subunit HflK PA4942 [imported] - Pseudomonas aeruginosa Length = 400 gb|AAG08327.1|AE004907_5 protease subunit HflK [Pseudomonas aeruginosa PAO1] Best-BlastP=> >nrprot 62% 1123.2
- Identities = 58/152 (38%), Positives = 87/152 (57%), Gaps = 8/152 (5%) gb|AAM09314.2| similar to Mus musculus Length = 243[Dictyostelium discoideum] (Mouse). Uridine-cytidine kinase 2 Best-BlastP=> >nrprot 36% 1124.2
 - [Ralstonia solanacearum] emb|CAD15343.1| PROBABLE ACETYLORNITHINE Best-BlastP=> >nrprot 68% Identities = 193/382 (50%), Positives = 265/382 (69%) ref[NP_519762.1| PROBABLE ACETYLORNITHINE Length = 397[Ralstonia solanacearum] DEACETYLASE (ACETYLORNITHINASE) PROTEIN DEACETYLASE (ACETYLORNITHINASE) PROTEIN 1125.2
 - Identities = 163/398 (40%), Positives = 239/398 (60%), Gaps = 2/398 (0%) ref[ZP_00033611.1| COG0477: Length = 470 [Burkholderia fungorum] Permeases of the major facilitator superfamily Best-BlastP=> >nrprot 55% 1126.3
- synthetase, putative [Pseudomonas putida KT2440] gb|AAN68679.1|AE016497_4 acetoacetyl-CoA synthetase, putative [Pseudomonas putida Best-BlastP=> >nrprot 72% Identities = 360/650 (55%), Positives = 472/650 (72%), Gaps = 4/650 (0%) ref[NP_745215.1] acetoacetyl-CoA Length = 6501127.3
- Best-BlastP=> >nrprot 50% Identities = 113/289 (39%), Positives = 170/289 (58%), Gaps = 13/289 (4%) refINP_904046.1 biotin synthesis protein [Chromobacterium violaceum ATCC 12472] gb|AAQ62035.1| biotin synthesis protein [Chromobacterium violaceum ATCC 12472] Length = 302 1128.3
- Best-BlastP=> >nrprot 57% Identities = 85/221 (38%), Positives = 135/221 (61%), Gaps = 12/221 (5%) ref[ZP_00086846.1| COG1040: Predicted Length = 246 fluorescens Pf0-1] amidophosphoribosyltransferases [Pseudomonas 1132.2
 - Best-BlastP=> >nrprot 50% Identities = 103/346 (29%), Positives = 172/346 (49%), Gaps = 13/346 (3%) ref|ZP_00082200.1| COG0750: Length = 355[Geobacter metallireducens] Predicted membrane-associated Zn-dependent proteases 1 1133.3
 - 1134.3 Best-BlastP=> >nrprot No Hits found
- Identities = 232/442 (52%), Positives = 307/442 (69%), Gaps = 1/442 (0%) ref[ZP_00043253.1| COG0790: FOG: PR repeat, SEL1 subfamily [Magnetococcus sp. MC-1] Best-BlastP=> >nrprot 62% 1136.3
- 1137.2 Best-BlastP=> >nrprot No Hits found
- 114.2 Best-BlastP=> >nrprot No Hits found
- Identities = 222/454 (48%), Positives = 312/454 (68%), Gaps = 4/454 (0%) ref[NP_820336.1] amino acid antiporter Length = 474 Coxiella burnetii RSA 493] gb/AAO90850.1 amino acid antiporter [Coxiella burnetii RSA 493] Best-BlastP=> >nrprot 65% 1142.4

- Best-BlastP=> >nrprot 64% Identities = 180/379 (47%), Positives = 254/379 (67%), Gaps = 4/379 (1%) ref[NP_800676.1| conserved hypothetical 2210633] splQ87GZ9|CLCA_VIBPA Voltage-gated CIC-type chloride channel clcA Length = 467dbj|BAC62509.1| conserved hypothetical protein [Vibrio parahaemolyticus] protein [Vibrio parahaemolyticus RIMD 1144.3
 - Best-BlastP=> >nrprot 72% Identities = 391/721 (54%), Positives = 522/721 (72%), Gaps = 11/721 (1%) ref[ZP_00110122.1| COG0512: Length = 734[Nostoc punctiforme] Anthranilate/para-aminobenzoate synthases component II 1145.3

1148.2

- 3est-BlastP=> >nrprot 56% Identities = 334/906 (36%), Positives = 512/906 (56%), Gaps = 35/906 (3%) ref[NP_289628.1| adenylylating enzyme Escherichia coli (strain O157:H7, Escherichia coli (strain 0157:H7, O157:H7 EDL933] O157:H7 EDL933] ref[NP_311963.1| glutamate-ammonia-ligase adenylyltransferase Length = 946 substrain EDL933) gblAAG58187.1|AE005534_9 adenylylating enzyme for glutamine synthetase [Escherichia coli O157:H7] pir[|H91120 glutamate-ammonia-ligase adenylyltransferase [imported] -0509952) pir||G85965 adenylylating enzyme for glutamine synthetase [imported] -0157:H7] dbj|BAB37359.1| glutamate-ammonia-ligase adenylyltransferase [Escherichia coli for glutamine synthetase [Escherichia coli substrain RIMD
 - 7120) plasmid Best-BlastP=> >nrprot 57% Identities = 53/167 (31%), Positives = 98/167 (58%), Gaps = 5/167 (2%) ref[NP_490158.1| probable Length = 169 acetyltransferase [Nostoc sp. PCC 7120] pir||AD2484 hypothetical protein alr7052 [imported] - Nostoc sp. (strain PCC pCC7120alpha dbj|BAB78136.1| ORF_ID:alr7052~probable acetyltransferase [Nostoc sp. PCC 7120] 115.2
 - Identities = 39/137 (28%), Positives = 67/137 (48%), Gaps = 10/137 (7%) ref[NP_824638.1] putative lipase Length = 286 Streptomyces avermitilis MA-4680] dbj|BAC71173.1| putative lipase [Streptomyces avermitilis MA-4680] Best-BlastP=> >nrprot 22% 1152.2
- membrane protein [Streptomyces coelicolor A3(2)] pir|[T35887 hypothetical protein SC9B10.18 Streptomyces coelicolor emb|CAA15808.1| Best-BlastP=> >nrprot 46% Identities = 36/136 (26%), Positives = 67/136 (49%), Gaps = 15/136 (11%) ref[NP_629974.1] putative integral Length = 312 putative integral membrane protein [Streptomyces coelicolor A3(2)] 1153.1
 - Best-BlastP=> >nrprot 48% Identities = 114/437 (26%), Positives = 213/437 (48%), Gaps = 37/437 (8%) ref[ZP_00107812.1| COG4325: Length = 449 Predicted membrane protein [Nostoc punctiforme] 1156.3
- Best-BlastP=> >nrprot 73% Identities = 550/943 (58%), Positives = 687/943 (72%), Gaps = 15/943 (1%) ref|ZP_00085403.1| COG0567: 2component, and related enzymes [Pseudomonas fluorescens oxoglutarate dehydrogenase complex, dehydrogenase (E1) Length = 943 1157.4
- Best-BlastP=> >nrprot 30% Identities = 61/269 (22%), Positives = 130/269 (48%), Gaps = 23/269 (8%) ref[NP_143635.1] chromosome assembly protein [Pyrococcus horikoshii] pir||F71190 probable chromosome assembly protein - Pyrococcus horikoshii dbj|BAA30917.1| 1179aa long Length = 1179 horikoshii] hypothetical chromosome assembly protein [Pyrococcus 1159.5
- Best-BlastP=> >nrprot 29% Identities = 42/123 (34%), Positives = 47/123 (38%), Gaps = 27/123 (21%) ref[NP_639328.1| hypothetical protein ATCC 33913] gb|AAM43210.1| hypothetical protein [Xanthomonas campestris pv Length = 131 Xanthomonas campestris pv. campestris str. ATCC 33913] campestris str. 116.1
- flexneri] ref[NP_858160.1| hypothetical protein [Shigella flexneri 2a] gb|AAK18345.1|AF348706_34 IS10 orf [Shigella flexneri] gb|AAL72480.1| Best-BlastP=> >nrprot 51% Identities = 136/392 (34%), Positives = 206/392 (52%), Gaps = 10/392 (2%) ref[NP_085189.1] IS10 orf [Shigella Length = 407 nypothetical protein [Shigella flexneri 2a] 1160.2

- Identities = 225/353 (63%), Positives = 269/353 (76%), Gaps = 2/353 (0%) ref[NP_756808.1| Uroporphyrinogen deċarboxylase [Escherichia coli CFT073] sp|Q8FB74|DCUP_ECOL6 Uroporphyrinogen decarboxylase (URO-D) (UPD) Length = 354gb|AAN83382.1|AE016770_182 Uroporphyrinogen decarboxylase [Escherichia coli CFT073] Best-BlastP=> >nrprot 75% 1161.2
- substrain RIMD 0509952) pir/|D85966 probable kinase ygiG [imported] Escherichia coli (strain O157:H7, Escherichia coli O157:H7 EDL933] ref|NP_311968.1 putative kinase [Escherichia coli O157:H7] pir||E91121 probable kinase [imported] Identities = 47/114 (41%), Positives = 71/114 (62%), Gaps = 2/114 (1%) ref[NP_289633.1| putative kinase substrain EDL933) gb|AAG58192.1|AE005535_4 putative kinase [Escherichia coli O157:H7 EDL933] dbj|BAB37364.1| putative kinase Length = 123Escherichia coli (strain 0157:H7, Best-BlastP=> >nrprot 62% Escherichia coli O157:H7] 1162.1
- Identities = 46/156 (29%), Positives = 68/156 (43%), Gaps = 10/156 (6%) ref[NP_747191.1| conserved hypothetical protein [Pseudomonas putida KT2440] gb|AAN70655.1|AE016709_7 conserved hypothetical protein [Pseudomonas putida KT2440] Best-BlastP=> >nrprot 29% 1163.2
- Best-BlastP=> >nrprot 61% Identities = 64/117 (54%), Positives = 83/117 (70%) ref[NP_755691.1| Hypothetical protein [Escherichia coli CFT073] Length = 365gb|AAN82265.1|AE016767_25 Hypothetical protein [Escherichia coli CFT073] 1168.2
- Best-BlastP=> >nrprot 99% Identities = 616/621 (99%), Positives = 618/621 (99%), Gaps = 1/621 (0%) gb/AAB09543.1| LprpoD 1169.3
- 117.1 Best-BlastP=> >nrprot No Hits found
- complement resistance protein precursor pir||C29835 TraTp protein Escherichia coli plasmid pED208 gb|AAA88375.1| traTp gene product Best-BlastP=> >nrprot 64% Identities = 104/241 (43%), Positives = 161/241 (66%), Gaps = 6/241 (2%) sp|P13980|TRT3_ECOLI TraT Length = 245gb|AAM90725.2| TraT [Salmonella typhi] 1171.3
- 172.2 Best-BlastP=> >nrprot No Hits found
- Best-BlastP=> >nrprot 74% Identities = 267/451 (59%), Positives = 334/451 (74%), Gaps = 9/451 (1%) ref|ZP_00065505.1| COG0486: Predicted Length = 456 GTPase [Microbulbifer degradans 2-40] 1174.2
 - 1176.5 Best-BlastP=> >nrprot No Hits found
- 1179.1 Best-BlastP=> >nrprot No Hits found
- 118.1 Best-BlastP=> >nrprot No Hits found
- Best-BlastP=> >nrprot 44% Identities = 66/232 (28%), Positives = 113/232 (48%), Gaps = 2/232 (0%) refINP_762946.1 AraC-type DNA-binding CMCP6] gb/AAO07936.1/AE016811_177 AraC-type DNA-binding domain-containing protein Length = 237domain-containing protein [Vibrio vulnificus CMCP61 Vibrio vulnificus 1180.2
- 3est-BlastP=> >nrprot 48% Identities = 88/274 (32%), Positives = 149/274 (54%), Gaps = 18/274 (6%) refINP_520074.1| PROBABLE FRANSMEMBRANE PROTEIN [Ralstonia solanacearum] emb|CAD15655.1| PROBABLE TRANSMEMBRANE PROTEIN [Ralstonia 1182.3
- 1183.4 Best-BlastP=> >nrprot No Hits found
- 1184.2

Best-BlastP=> >nrprot 54% Identities = 133/395 (33%), Positives = 216/395 (54%), Gaps = 6/395 (1%) refINP_618268.1| amino acid transporter Methanosarcina acetivorans str. C2A] gb/AAM06748.1| amino acid transporter [Methanosarcina acetivorans str. C2A]

- Best-BlastP=> >nrprot 31% Identities = 52/171 (30%), Positives = 82/171 (47%), Gaps = 7/171 (4%) ref|NP_149698.1| 235L [Invertebrate Length = 265 ridescent virus 6] gb|AAK82096.1|AF303741_235 235L [Chilo iridescent virus] 1186.2
- Best-BlastP=> >nrprot 99% Identities = 737/749 (98%), Positives = 746/749 (99%) sp|Q9WXB9|CATA_LEGPN Peroxidase/catalase (Catalaseperoxidase) dbj|BAA78342.1| catalase-peroxidase [Legionella pneumophila] gb|AAG37106.1|AF276752_1 catalase-peroxidase [Legionella Length = 749pneumophilal 1188.2
 - Best-BlastP=> >nrprot 51% Identities = 53/124 (42%), Positives = 77/124 (62%), Gaps = 2/124 (1%) dbj|BAA75251.1| Similar to IS1301 of Length = 255actinomycetemcomitans] Veisseria meningitidis [Actinobacillus 119.1
 - Best-BlastP=> >nrprot 56% Identities = 256/709 (36%), Positives = 406/709 (57%), Gaps = 25/709 (3%) ref|ZP_00065821.1| COG1249 related enzymes dihydrolipoamide dehydrogenase (E3) component, and Length = 704 Pyruvate/2-oxoglutarate dehydrogenase complex, Microbulbifer degradans 2-40] 1190.4
 - 1193.2 Best-BlastP=> >nrprot No Hits found
- str. ATCC 33913] gb/AAM40726.1| glutaryl-CoA dehydrogenase [Xanthomonas Best-BlastP=> >nrprot 72% Identities = 223/383 (58%), Positives = 278/383 (72%), Gaps = 1/383 (0%) refINP_636802.1| glutaryl-CoA Length = 387 dehydrogenase [Xanthomonas campestris pv. campestris str. ATCC 33913] campestris pv. campestris 1194.3
 - Identities = 73/110 (66%), Positives = 90/110 (81%) ref|ZP_00067987.1| COG0091: Ribosomal protein L22 Length = 110 Microbulbifer degradans 2-40] Best-BlastP=> >nrprot 80% 1199.4
- [Coxiella burnetii RSA 493] sp|O85388|RS3_COXBU 30S ribosomal protein S3 gb|AAO89802.1| ribosomal protein S3 [Coxiella burnetii RSA 493] Best-BlastP=> >nrprot 82% Identities = 150/220 (68%), Positives = 180/220 (81%), Gaps = 1/220 (0%) ref|NP_819288.1| ribosomal protein S3 -ength = 227 1201.1
- Best-BlastP=> >nrprot 89% Identities = 113/137 (82%), Positives = 124/137 (90%) refINP_742627.1| ribosomal protein L16 [Pseudomonas Length = 137putida KT2440] gb|AAN66091.1|AE016238_9 ribosomal protein L16 [Pseudomonas putida KT2440] 1202.2
 - Best-BlastP=> >nrprot 16% Identities = 428/1182 (36%), Positives = 630/1182 (53%), Gaps = 81/1182 (6%) ref|ZP_00108802.1| hypothetical Length = 2315protein [Nostoc punctiforme] 1205.5
 - 1207.2 Best-BlastP=> >nrprot No Hits found
- Best-BlastP=> >nrprot 42% Identities = 45/124 (36%), Positives = 55/124 (44%), Gaps = 4/124 (3%) pir||F71456 hypothetical protein PH0308 Length = 215Pyrococcus horikoshii dbj|BAA29381.1| 215aa long hypothetical protein [Pyrococcus horikoshii] 1208.1
 - Best-BlastP=> >nrprot 84% Identities = 346/487 (71%), Positives = 417/487 (85%), Gaps = 1/487 (0%) ref|ZP_00087773.1| COG0516: IMP Length = 506Pf0-1] dehydrogenase/GMP reductase [Pseudomonas fluorescens 1209.2
 - Best-BlastP=> >nrprot 59% Identities = 58/101 (57%), Positives = 67/101 (66%), Gaps = 1/101 (0%) ref[ZP_00091135.1| COG2852 Length = 150vinelandii] Uncharacterized protein conserved in bacteria [Azotobacter 121.1
- gb|AAN44053.1|AE015270_8 GMP synthetase [Shigella flexneri 2a str. 301] gb|AAP17880.1| GMP synthetase (glutamine-hydrolyzing) [Shigella Best-BlastP=> >nrprot 81% Identities = 354/523 (67%), Positives = 427/523 (81%), Gaps = 1/523 (0%) refINP_708346.1| GMP synthetase [Shigella flexneri 2a str. 301] ref|NP_838070.1| GMP synthetase (glutamine-hydrolyzing) [Shigella flexneri 2a str. 1210.2

- TTO1] emb|CAE14921.1| unnamed protein product [Photorhabdus luminescens subsp. Identities = 47/90 (52%), Positives = 69/90 (76%), Gaps = 1/90 (1%) ref[NP_929783.1| hypothetical protein Photorhabdus luminescens subsp. laumondii Length = 94 Best-BlastP=> >nrprot 19% 1211.3
- Best-BlastP=> >nrprot 65% Identities = 193/394 (48%), Positives = 248/394 (62%), Gaps = 22/394 (5%) ref[NP_249543.1| chitin-binding protein CbpD precursor [Pseudomonas aeruginosa PA01] pir[[F83538 chitin-binding protein CbpD precursor PA0852 [imported] Pseudomonas aeruginosa (strain PAO1) gb/AAG04241.1/AE004520_4 chitin-binding protein CbpD precursor [Pseudomonas aeruginosa PAO1] 1213.2
- HYPOTHETICAL PROTEIN [Sinorhizobium meliloti] emb[CAC47611.1] CONSERVED HYPOTHETICAL PROTEIN [Sinorhizobium meliloti] Identities = 172/363 (47%), Positives = 233/363 (64%), Gaps = 5/363 (1%) ref[NP_387138.1| CONSERVED Best-BlastP=> >nrprot 64% 1215.3
- vartitioning protein [Xylella fastidiosa Temecula1] splQ87BY1|PARB_XYLFT Probable chromosome partitioning protein parB gb|AAO29164.1| Best-BlastP=> >nrprot 69% Identities = 90/159 (56%), Positives = 115/159 (72%), Gaps = 4/159 (2%) refINP_779515.1| chromosome Length = 310chromosome partitioning protein [Xylella fastidiosa Temecula1] 122.3
- Best-BlastP=> >nrprot 77% Identities = 332/507 (65%), Positives = 395/507 (77%), Gaps = 3/507 (0%) dbj|BAB19801.1| piperideine-6-Length = 510 lutescens carboxylate dehydrogenase ['Flavobacterium' 1220.3
 - 1222.5 Best-BlastP=> >nrprot No Hits found
- Best-BlastP=> >nrprot 39% Identities = 73/287 (25%), Positives = 119/287 (41%), Gaps = 38/287 (13%) refINP_521275.1| PROBABLE SIGNAL PEPTIDE PROTEIN [Ralstonia solanacearum] emb|CAD16942.1| PROBABLE SIGNAL PEPTIDE PROTEIN [Ralstonia solanacearum] Length = 278 1223.2
- Identities = 212/266 (79%), Positives = 237/266 (89%) emb[CAC35728.1] OXA-29 [Fluoribacter gormanii] Best-BlastP=> >nrprot 88% 1224.2
- AF2122/97] emb[CAD94271.1] CONSERVED HYPOTHETICAL PROTEIN Best-BlastP=> >nrprot 40% Identities = 90/292 (30%), Positives = 143/292 (48%), Gaps = 26/292 (8%) ref[NP 855062.1| CONSERVED Length = 439 HYPOTHETICAL PROTEIN [Mycobacterium bovis subsp. bovis AF2122/97] Mycobacterium bovis subsp. bovis 1225.2
- Best-BlastP=> >nrprot 55% Identities = 129/350 (36%), Positives = 190/350 (54%), Gaps = 22/350 (6%) ref[NP_629326.1| putative sulfurylase Length = 392 Streptomyces coelicolor A3(2)] emb[CAC01308.1] putative sulfurylase [Streptomyces coelicolor A3(2)] 1226.2
 - Best-BlastP=> >nrprot 48% Identities = 66/202 (32%), Positives = 109/202 (53%), Gaps = 10/202 (4%) ref|ZP_00107215.1| COG0637: Predicted Length = 242 punctiforme] phosphatase/phosphohexomutase [Nostoc 1227.1
 - 1229.3 Best-BlastP=> >nrprot No Hits found
- Best-BlastP=> >nrprot 59% Identities = 201/462 (43%), Positives = 284/462 (61%), Gaps = 28/462 (6%) ref|NP_253634.1| N-acetylmuramoyl-Lalanine amidase [Pseudomonas aeruginosa PA01] pir||G83028 N-acetylmuramoyl-L-alanine amidase PA4947 [imported] - Pseudomonas aeruginosa (strain PAO1) gb|AAG08332.1|AE004907_10 N-acetylmuramoyl-L-alanine amidase [Pseudomonas aeruginosa PAO1] 123.2
- Best-BlastP=> >nrprot 81% Identities = 364/516 (70%), Positives = 429/516 (83%) ref[NP_819831.1] peptide chain release factor 3 [Coxiella burnetii RSA 493] gb/AAO90345.1| peptide chain release factor 3 [Coxiella burnetii RSA 493] 1230.2
 - 1231.3 Best-BlastP=> >nrprot No Hits found

- Identities = 44/137 (32%), Positives = 70/137 (51%), Gaps = 1/137 (0%) ref[NP_800044.1] putative acetyltransferase Length = 140[Vibrio parahaemolyticus RIMD 2210633] dbj|BAC61877.1| putative acetyltransferase [Vibrio parahaemolyticus] Best-BlastP=> >nrprot 28% 1235.3
- Identities = 172/176 (97%), Positives = 174/176 (98%) gb|AAM00633.1| unknown [Legionella pneumophila] Best-BlastP=> >nrprot 65% Length = 176 1236.2
- Identities = 47/132 (35%), Positives = 73/132 (55%), Gaps = 4/132 (3%) ref[NP_832446.1| Acetyltransferase Length = 141 Bacillus cereus ATCC 14579] gb|AAP09647.1| Acetyltransferase [Bacillus cereus ATCC 14579] Best-BlastP=> >nrprot 50% 1237.2
- 1238.2 Best-BlastP=> >nrprot No Hits found
- 1239.2 Best-BlastP=> >nrprot No Hits found
- Identities = 66/144 (45%), Positives = 95/144 (65%) refINP_716232.1| conserved hypothetical protein TIGR00150 MR-1] gb/AAN53677.1/AE015507_3 conserved hypothetical protein TIGR00150 [Shewanella oneidensis Best-BlastP=> >nrprot 63% Length = 152 Shewanella oneidensis 124.1
 - 1242.2

Identities = 138/191 (72%), Positives = 160/191 (83%) refINP_353171.1| AGR_C_216p [Agrobacterium tumefaciens] C58 (U. Washington)] splQ8UJ06|UPP_AGRT5 Uracil Cereon) pir||Al2592 uracil phosphoribosyltransferase [imported] phosphoribosyltransferase (UMP pyrophosphorylase) (UPRTase) pir||C97375 uracil phosphoribosyltransferase (UMP pyrophosphorylase) tumefaciens (strain C58, Dupont) gb|AAK85956.1| AGR_C_216p [Agrobacterium tumefaciens str. C58 (Cereon)] C58 (U. Washington)] refINP_530843.1| uracil phosphoribosyltransferase [Agrobacterium tumefaciens str. gb[AAL41159.1] uracil phosphoribosyltransferase [Agrobacterium tumefaciens str. [imported] - Agrobacterium tumefaciens (strain C58, Best-BlastP=> >nrprot 82% Agrobacterium

- 1244.2 Best-BlastP=> >nrprot No Hits found
- 1245.1 Best-BlastP=> >nrprot No Hits found
- Best-BlastP=> >nrprot 22% Identities = 82/298 (27%), Positives = 130/298 (43%), Gaps = 56/298 (18%) pir||T09051 PepA protein -Pseudomonas aeruginosa gb|AAC16023.1| ExoU [Pseudomonas aeruginosa] gb|AAC38269.1| PepA [Pseudomonas aeruginosa] Length = 687 gb|AAP82959.1| type III effector protein [Pseudomonas aeruginosa] 1246.2
- Best-BlastP=> >nrprot 30% Identities = 37/119 (31%), Positives = 57/119 (47%), Gaps = 9/119 (7%) emb|CAD21525.1| hypothetical protein Length = 155 Taenia solium] 1249.2
- 125.2
- Best-BlastP=> >nrprot 59% Identities = 227/486 (46%), Positives = 293/486 (60%), Gaps = 4/486 (0%) ref|NP_820087.1| conserved hypothetical Best-BlastP=> >nrprot 64% Identities = 74/160 (46%), Positives = 105/160 (65%), Gaps = 1/160 (0%) ref[NP_820968.1] dihydrofolate reductase protein [Coxiella burnetii RSA 493] gb|AAO90601.1| conserved hypothetical protein [Coxiella burnetii RSA 493] 1250.1
 - aeruginosa PA01] sp[Q9I5U4[PXA1_PSEAE 4-hydroxythreonine-4-phosphate dehydrogenase 1 Best-BlastP=> >nrprot 68% Identities = 176/321 (54%), Positives = 222/321 (69%), Gaps = 4/321 (1%) refINP_249284.1| pyridoxal phosphate Pseudomonas aeruginosa (strain PAO1) gb/AAG03982.1/AE004495_6 pyridoxal phosphate biosynthetic protein PdxA [Pseudomonas (4-(phosphohydroxy)-L-threonine dehydrogenase 1) pir||A83572 pyridoxal phosphate biosynthetic protein PdxA PA0593 [imported] Length = 161 Coxiella burnetii RSA 493] gb/AAO91482.1 dihydrofolate reductase [Coxiella burnetii RSA 493] biosynthetic protein PdxA [Pseudomonas Length = 328aeruginosa PAO1] 1251.2

Identities = 242/415 (58%), Positives = 300/415 (72%), Gaps = 5/415 (1%) refINP_820851.1| conserved hypothetical Length = 435 protein [Coxiella burnetii RSA 493] gb|AAO91365.1| conserved hypothetical protein [Coxiella burnetii RSA 493] Best-BlastP=> >nrprot 69%

1257.5

- Identities = 175/312 (56%), Positives = 217/312 (69%), Gaps = 2/312 (0%) ref[ZP_00125152.1| COG0189: enzyme (glutaminyl transferase) [Pseudomonas syringae Glutathione synthase/Ribosomal protein S6 modification Best-BlastP=> >nrprot 67% Length = 3191258.1
- Identities = 470/475 (98%), Positives = 475/475 (100%) emb|CAD42896.1| flagellin [Legionella pneumophila Length = 475 Best-BlastP=> >nrprot 99% serogroup 1] 126.2
- solanacearum] emb|CAD16666.1| PROBABLE LIPOPROTEIN Identities = 92/211 (43%), Positives = 126/211 (59%), Gaps = 3/211 (1%) refINP_521080.1| PROBABLE Length = 269solanacearum] -IPOPROTEIN PRECURSOR (VACJ) TRANSMEMBRANE [Ralstonia PRECURSOR (VACJ) TRANSMEMBRANE [Ralstonia Best-BlastP=> >nrprot 48% 1260.3
- Identities = 286/564 (50%), Positives = 370/564 (65%), Gaps = 8/564 (1%) gb|AAB16855.1| pyruvate decarboxylase Length = 607Best-BlastP=> >nrprot 66% Arabidopsis thaliana] 1261.2
- aminopeptidase [Chromobacterium violaceum ATCC 12472] gb[AAQ57736.1] probable aminopeptidase [Chromobacterium violaceum ATCC Identities = 97/291 (33%), Positives = 159/291 (54%), Gaps = 16/291 (5%) ref|NP 899726.1| probable Best-BlastP=> >nrprot 39% Length = 415 1264.3
- Identities = 119/262 (45%), Positives = 162/262 (61%), Gaps = 6/262 (2%) ref|ZP_00051284.1| COG0656: Aldo/keto Length = 291[Magnetospirillum magnetotacticum] reductases, related to diketogulonate reductase Best-BlastP=> >nrprot 58% 1265.5
 - Identities = 38/108 (35%), Positives = 55/108 (50%), Gaps = 8/108 (7%) ref[NP_616727.1| conserved hypothetical C2Aj gb/AAM05207.1| conserved hypothetical protein [Methanosarcina acetivorans str protein [Methanosarcina acetivorans str. Best-BlastP=> >nrprot 20% Length = 266 1267.3
- 1268.2
- Length = 310Best-BlastP=> >nrprot 37% Identities = 32/94 (34%), Positives = 48/94 (51%), Gaps = 6/94 (6%) sp|Q02910|CPN_DROME Calphotin Identities = 82/293 (27%), Positives = 144/293 (49%), Gaps = 33/293 (11%) ref[XP_313252.1] ENSANGP00000010487 [Anopheles gambiae] gb|EAA08759.1| ENSANGP00000010487 [Anopheles gambiae str. PEST] Best-BlastP=> >nrprot 46% 127.4
- Best-BlastP=> >nrprot 70% Identities = 73/144 (50%), Positives = 102/144 (70%), Gaps = 2/144 (1%) ref[NP_241923.1] BH1057~unknown melanogaster) gb[AAA28405.1] calcium-binding protein pir||A47282 calcium-binding protein calphotin - fruit fly (Drosophila 1270.2
 - conserved protein [Bacillus halodurans] sp|Q9RC41|YA57_BACHD Hypothetical protein BH1057 pir||A83782 hypothetical protein BH1057 (strain C-125) dbj|BAA83958.1| YHDE [Bacillus halodurans] dbj|BAB04776.1| BH1057~unknown Length = 146 conserved protein [Bacillus halodurans] [imported] - Bacillus halodurans
- Identities = 35/70 (50%), Positives = 53/70 (75%) splP17724|GLB_TETPY Myoglobin (Hemoglobin) pir||A36270 nemoglobin - Tetrahymena pyriformis dbj|BAA03015.1| hemoglobin [Tetrahymena pyriformis] Best-BlastP=> >nrprot 61% 1271.3

Identities = 89/318 (27%), Positives = 147/318 (46%), Gaps = 13/318 (4%) ref[NP_253021.1] probable ferredoxin Length = 308reductase [Pseudomonas aeruginosa PA01] pir||G83104 probable ferredoxin reductase PA4331 [imported] - Pseudomonas strain PAO1) gb[AAG07719.1|AE004849_6 probable ferredoxin reductase [Pseudomonas aeruginosa PAO1] 3est-BlastP=> >nrprot 46%

1272.4

- Identities = 54/200 (27%), Positives = 95/200 (47%), Gaps = 8/200 (4%) gb/AAM88782.1 hypothetical protein Length = 247 Best-BlastP=> >nrprot 40% Photorhabdus luminescens] 1273.4
- Identities = 107/291 (36%), Positives = 171/291 (58%), Gaps = 1/291 (0%) gb/AAM88781.1| MhpE-like protein Length = 312 Best-BlastP=> >nrprot 57% Photorhabdus luminescens] 1275.2
- tengcongensis] gb[AAM25889.1| UDP-N-acetylmuramyl tripeptide synthase Identities = 100/294 (34%), Positives = 162/294 (55%), Gaps = 11/294 (3%) ref[NP_624285.1| UDP-N-Length = 879 acetylmuramyl tripeptide synthase [Thermoanaerobacter tengcongensis] Best-BlastP=> >nrprot 48% Thermoanaerobacter 1277.2
- Identities = 104/386 (26%), Positives = 176/386 (45%), Gaps = 49/386 (12%) gb[AAN83921.1] hypothetical protein Length = 427Best-BlastP=> >nrprot 43% Aplysia californica] 1278.2
- nfluenzae Rd] sp|P44255|YFCM_HAEIN Hypothetical protein HI1563 pir||D64036 hypothetical protein HI1563 Haemophilus influenzae (strain Best-BlastP=> >nrprot 76% Identities = 107/173 (61%), Positives = 136/173 (78%) ref[NP_439712.1| hypothetical protein [Haemophilus Length = 178 KW20) gb[AAC23212.1] conserved hypothetical protein [Haemophilus influenzae Rd] 1279.3
 - Length Best-BlastP=> >nrprot 15% Identities = 110/113 (97%), Positives = 110/113 (97%) gb|AAO61471.1| LidA [Legionella pneumophila] 1280.3
- Best-BlastP=> >nrprot 44% Identities = 155/493 (31%), Positives = 230/493 (46%), Gaps = 37/493 (7%) gb[AAC35592.1| LphB [Legionella Length = 518 pneumophila] 1282.3
- Identities = 180/429 (41%), Positives = 260/429 (60%), Gaps = 9/429 (2%) refINP_716339.1 conserved hypothetical protein [Shewanella oneidensis MR-1] gb|AAN53784.1|AE015516_6 conserved hypothetical protein [Shewanella oneidensis MR-1] Best-BlastP=> >nrprot 59% 1283.2
- Identities = 25/75 (33%), Positives = 43/75 (57%) refINP_796782.1| hypothetical protein VP0403 [Vibrio Length = 111 2210633] dbj|BAC58666.1| hypothetical protein [Vibrio parahaemolyticus] Best-BlastP=> >nrprot 45% parahaemolyticus RIMD 1284.2
- Identities = 71/260 (27%), Positives = 123/260 (47%), Gaps = 17/260 (6%) gb|AAK19884.1| putative methoxymalonyl Length = 863 CoA synthase [Polyangium cellulosum] Best-BlastP=> >nrprot 19% 1286.3
- Identities = 278/933 (29%), Positives = 458/933 (49%), Gaps = 56/933 (6%) ref[ZP_00089642.1| hypothetical protein Length = 973 Best-BlastP=> >nrprot 49% Azotobacter vinelandii] 1287.3 1288.2
- Identities = 39/130 (30%), Positives = 67/130 (51%), Gaps = 8/130 (6%) ref[XP_314825.1] ENSANGP0000011098 Length = 1842 Anopheles gambiae] gb|EAA10144.1| ENSANGP00000011098 [Anopheles gambiae str. PEST] Best-BlastP=> >nrprot No Hits found Best-BlastP=> >nrprot 14%
- Best-BlastP=> >nrprot 62% Identities = 91/182 (50%), Positives = 123/182 (67%), Gaps = 2/182 (1%) ref|ZP_00013245.1| COG2353: Uncharacterized conserved protein [Rhodospirillum rubrum] 1293.4

- Best-BlastP=> >nrprot 59% Identities = 77/177 (43%), Positives = 105/177 (59%), Gaps = 7/177 (3%) refINP_902948.1| probable cytochrome b561 [Chromobacterium violaceum ATCC 12472] gb|AAQ60942.1| probable cytochrome b561 [Chromobacterium violaceum ATCC 12472] 1294.2
- Best-BlastP=> $\frac{47\%}{600}$ Identities = $\frac{64}{169}$ (37%), Positives = $\frac{91}{169}$ (53%), Gaps = $\frac{1}{169}$ (0%) ref[NP_422185.1] conserved hypothetical protein [Caulobacter crescentus CB15] pir||E87669 conserved hypothetical protein CC3391 [imported] - Caulobacter gb|AAK25353.1| conserved hypothetical protein [Caulobacter crescentus CB15] 1295.1
 - Best-BlastP=> >nrprot 53% Identities = 150/437 (34%), Positives = 246/437 (56%), Gaps = 6/437 (1%) gb|AAD28727.1|AF112468_6 TraH Length = 427 protein precursor [Salmonella typhimurium] 1299.3

Length = 457

- Best-BlastP=> >nrprot 40% Identities = 57/260 (21%), Positives = 111/260 (42%), Gaps = 28/260 (10%) refINP_125735.1| hypothetical protein [Pyrococcus abyssi] pir||G75189 hypothetical protein PAB2321 - Pyrococcus abyssi (strain Orsay) emb|CAB48966.1| Hypothetical protein Length = 249Pyrococcus abyssi] 13.1
 - Best-BlastP=> >nrprot 50% Identities = 184/554 (33%), Positives = 274/554 (49%), Gaps = 94/554 (16%) refINP_841634.1| Flagellar hook-19718] emb|CAD85506.1| Flagellar hook-associated protein 2 [Nitrosomonas associated protein 2 [Nitrosomonas europaea ATCC Length = 481 19718] europaea ATCC 130.4
 - Best-BlastP=> >nrprot 54% Identities = 129/348 (37%), Positives = 197/348 (56%), Gaps = 10/348 (2%) ref|ZP_00130398.1| COG2200: FOG: Length = 367EAL domain [Desulfovibrio desulfuricans G20] 1302.2
 - Identities = 168/252 (66%), Positives = 207/252 (82%) ref[ZP_00125838.1] COG0024: Methionine aminopeptidase Length = 260syringae B728al Best-BlastP=> >nrprot 80% Pseudomonas syringae pv. 1303.3
- | Identities = 1033/1066 (96%), Positives = 1049/1066 (98%) gb|AAM00612.1| chemiosmotic efflux system protein A-Length = 1066 pneumophila] Best-BlastP=> >nrprot 98% ike protein [Legionella 1306.5
 - Identities = 389/418 (93%), Positives = 407/418 (97%) gb|AAM00611.1| proline/glycine betaine transporter-like Length = 422 pneumophila] Best-BlastP=> >nrprot 97% protein [Legionella 307.4
- CV2125 [Chromobacterium violaceum ATCC 12472] gb|AAQ59798.1| hypothetical protein CV2125 [Chromobacterium violaceum ATCC 12472] Best-BlastP=> >nrprot 26% Identities = 26/88 (29%), Positives = 42/88 (47%), Gaps = 15/88 (17%) ref|NP_901795.1| hypothetical protein 131.2
- Best-BlastP=> >nrprot 57% Identities = 170/474 (35%), Positives = 278/474 (58%), Gaps = 14/474 (2%) ref|ZP_00129665.1| COG1538: Outer Length = 494 membrane protein [Desulfovibrio desulfuricans G20] 1314.3
 - Best-BlastP=> >nrprot No Hits found 1317.3
 - 1318.4

Best-BlastP=> >nrprot 64% Identities = 110/230 (47%), Positives = 162/230 (70%), Gaps = 10/230 (4%) ref|NP_819966.1| conserved hypothetical protein [Coxiella burnetii RSA 493] gb|AAO90480.1| conserved hypothetical protein [Coxiella burnetii RSA 493]

- Best-BlastP=> >nrprot 67% Identities = 365/741 (49%), Positives = 498/741 (67%), Gaps = 11/741 (1%) ref|ZP_00010561.1| COG0068: Length = 772 Hydrogenase maturation factor [Rhodopseudomonas palustris] 1320.3
- typhimurium LT2] gb/AAL22745.1| putative MFS family tranport protein [Salmonella typhimurium Best-BlastP=> >nrprot 60% Identities = 173/452 (38%), Positives = 280/452 (61%), Gaps = 6/452 (1%) ref|NP_462786.1| putative MFS family tranport protein (1st mdule) [Salmonella 1322.2

Best-BlastP=> >nrprot No Hits found

Best-BlastP=> >nrprot 28% Identities = 76/274 (27%), Positives = 130/274 (47%), Gaps = 35/274 (12%) gb|EAA21537.1| Plasmodium Length = 565alciparum CDPK2 protein [Plasmodium yoelii yoelii] 1324.4

Identities = 311/589 (52%), Positives = 419/589 (71%) refINP_820983.1| arginyl-tRNA synthetase [Coxiella burnetii Best-BlastP=> >nrprot 71% 1325.2

Best-BlastP=> >nrprot 55% Identities = 121/344 (35%), Positives = 196/344 (56%), Gaps = 8/344 (2%) ref|NP_819936.1| transporter, putative Length = 592RSA 493] gb|AAO91497.1| arginyl-tRNA synthetase [Coxiella burnetii RSA 493] 1328.2

Length = 376 [Coxiella burnetii RSA 493] gb|AAO90450.1| transporter, putative [Coxiella burnetii RSA 493] 1330.5

Best-BlastP=> >nrprot 69% Identities = 372/682 (54%), Positives = 478/682 (70%), Gaps = 4/682 (0%) ref|ZP_00092220.1| COG1200: RecG-Length = 1006 1331.3

Best-BlastP=> >nrprot 55% Identities = 125/409 (30%), Positives = 223/409 (54%), Gaps = 18/409 (4%) ref|NP_391052.1| alternate gene name: subtilis gb|AAA22318.1| B competence protein emb|CAB07900.1| unknown [Bacillus subtilis] emb|CAB15162.1| yuxH [Bacillus subtilis subsp. comB, yufA [Bacillus subtilis] sp|P14203|YUXH_BACSU Hypothetical protein yuxH pir||BVBSCB competence protein ComB (yuxH) - Bacillus Length = 409

Best-BlastP=> >nrprot 77% Identities = 308/477 (64%), Positives = 377/477 (79%) ref[NP_840694.1| Glycine cleavage system P-protein 19718] emb|CAD84521.1| Glycine cleavage system P-protein [Nitrosomonas europaea ATCC Nitrosomonas europaea ATCC Length = 483 19718] 1332.3

Best-BlastP=> >nrprot No Hits found

Best-BlastP=> >nrprot No Hits found 1335.2

Identities = 365/371 (98%), Positives = 369/371 (99%) gb|AAM00605.1| florfenicol efflux pump-like protein Best-BlastP=> >nrprot 99% 1336.3

Length = 371 Legionella pneumophila] 1338.3 Best-BlastP=> >nrprot 76% Identities = 406/673 (60%), Positives = 510/673 (75%), Gaps = 8/673 (1%) ref|NP_837722.1| methionine tRNA Best-BlastP=> >nrprot 79% Identities = 109/185 (58%), Positives = 151/185 (81%) ref|NP_770170.1| blr3530 [Bradyrhizobium japonicum] synthetase [Shigella flexneri 2a str. 2457T] gb|AAP17531.1| methionine tRNA synthetase [Shigella flexneri 2a str. 2457T] 1339.2

Length = 206dbj|BAC48795.1| blr3530 [Bradyrhizobium japonicum USDA 110]

Best-BlastP=> >nrprot No Hits found 1341.2

Best-BlastP=> >nrprot 49% Identities = 73/251 (29%), Positives = 130/251 (51%), Gaps = 16/251 (6%) gb|AAM90720.1| TraF [Salmonella typhi] 1342.3

Best-BlastP=> >nrprot 14% Identities = 30/113 (26%), Positives = 51/113 (45%), Gaps = 15/113 (13%) ref|ZP_00101173.1| hypothetical protein Length = 367 Desulfitobacterium hafniense] 1344.3 1345.3 Best-BlastP=> >nrprot 65% Identities = 173/366 (47%), Positives = 239/366 (65%), Gaps = 7/366 (1%) ref|NP_778380.1| conserved hypothetical 19718] emb|CAD85427.1| Uncharacterized protein family UPF0006 [Nitrosomonas Best-BlastP=> >nrprot 69% Identities = 135/258 (52%), Positives = 183/258 (70%), Gaps = 5/258 (1%) ref|NP_841557.1| Uncharacterized protein [Xylella fastidiosa Temecula1] gb[AAO28029.1| conserved hypothetical protein [Xylella fastidiosa Temecula1] protein family UPF0006 [Nitrosomonas europaea ATCC europaea ATCC 1346.3

Identities = 82/333 (24%), Positives = 152/333 (45%), Gaps = 43/333 (12%) ref[NP_819818.1] multidrug resistance Length = 331protein [Coxiella burnetii RSA 493] gb|AAO90332.1| multidrug resistance protein [Coxiella burnetii RSA 493] 3est-BlastP=> >nrprot 43%

1354.2 Best-BlastP=> >nrprot No Hits found

1353.3

[Enterococcus faecalis V583] gb|AAO80578.1| amino acid ABC transporter, amino acid-Identities = 72/242 (29%), Positives = 117/242 (48%), Gaps = 8/242 (3%) ref[NP_814508.1] amino acid ABC Length = 722 [Enterococcus faecalis V583] ransporter, amino acid-binding/permease protein Best-BlastP=> >nrprot 44% binding/permease protein 1356.2

1357.2

Identities = 90/146 (61%), Positives = 107/146 (73%) refINP_718466.1| Yail/YqxD family protein [Shewanella Length = 151 oneidensis MR-1] gb|AAN55910.1|AE015727_10 Yail/YqxD family protein [Shewanella oneidensis MR-1] Best-BlastP=> >nrprot 71%

Identities = 357/357 (100%), Positives = 357/357 (100%) emb[CAD43479.1] O-acetyltransferase [Legionella Length = 357 Best-BlastP=> >nrprot 99% 1359.2

Identities = 112/443 (25%), Positives = 195/443 (44%), Gaps = 75/443 (16%) ref[NP_845547.1| conserved nypothetical protein [Bacillus anthracis str. Ames] gb|AAP27033.1| conserved hypothetical protein [Bacillus anthracis str. Ames] Best-BlastP=> >nrprot 39% 1360.6

Identities = 99/312 (31%), Positives = 162/312 (51%), Gaps = 25/312 (8%) ref[NP_250993.1| hypothetical protein Pseudomonas aeruginosa PA01] pir||A83358 hypothetical protein PA2303 [imported] - Pseudomonas aeruginosa Length = 339 gb|AAG05691.1|AE004656_3 hypothetical protein PA2303 [Pseudomonas aeruginosa PAO1] Best-BlastP=> >nrprot 46% 1361.3

Identities = 570/607 (93%), Positives = 588/607 (96%) gb|AAK00285.1|AF288536_7 unknown [Legionella Length = 607Best-BlastP=> >nrprot 96% ongbeachae 1362.3

Identities = 505/534 (94%), Positives = 516/534 (96%) gb/AAK00286.1 AF288536_8 possible sensor kinase protein Length = 534 Best-BlastP=> >nrprot 96% Legionella longbeachae] 1364.2

1366.3 Best-BlastP=> >nrprot No Hits found

Identities = 55/123 (44%), Positives = 83/123 (67%) ref[ZP_00047813.1| COG2391: Predicted transporter magnetotacticum] Length = 155 magnetotacticum] component [Magnetospirillum Best-BlastP=> >nrprot 58% 1367.3

1368.2 Best-BlastP=> >nrprot No Hits found

1369.2

Identities = 58/133 (43%), Positives = 92/133 (69%) ref[NP_800456.1] conserved hypothetical protein [Vibrio Length = 139 2210633] dbj|BAC62289.1| conserved hypothetical protein [Vibrio parahaemolyticus] Best-BlastP=> >nrprot 63% parahaemolyticus RIMD

Identities = 66/135 (48%), Positives = 87/135 (64%), Gaps = 2/135 (1%) ref[NP_819361.1] conserved domain protein Length = 214 Coxiella burnetii RSA 493] gb|AAO89875.1| conserved domain protein [Coxiella burnetii RSA 493] Best-BlastP=> >nrprot 56% 1370.5

1372.2 Best-BlastP=> >nrprot No Hits found

1373.1 Best-BlastP=> >nrprot No Hits found

Best-BlastP=> >nrprot 29% Identities = 51/102 (50%), Positives = 69/102 (67%) ref[NP_790719.1] conserved hypothetical protein [Pseudomonas str. DC3000] gb/AAO54414.1| conserved hypothetical protein [Pseudomonas syringae pv. tomato syringae pv. tomato Length = 249 1375.2

- Identities = 38/97 (39%), Positives = 53/97 (54%), Gaps = 3/97 (3%) ref[ZP_00030779.1| hypothetical protein Length = 182 Best-BlastP=> >nrprot 40% Burkholderia fungorum] 1376.2
 - Best-BlastP=> >nrprot No Hits found 1377.5
- Best-BlastP=> >nrprot No Hits found 1378.5
- Best-BlastP=> >nrprot 15% Identities = 51/177 (28%), Positives = 81/177 (45%), Gaps = 26/177 (14%) ref|ZP_00071821.1| COG3240: 1380.3
- Identities = 61/216 (28%), Positives = 98/216 (45%), Gaps = 23/216 (10%) gb|AAK00282.1|AF288536_4 unknown Length = 418 erythraeum IMS101] Phospholipase/lecithinase/hemolysin [Trichodesmium Best-BlastP=> >nrprot 40% 1385.2
- Identities = 56/108 (51%), Positives = 72/108 (66%) gb|AAL17787.1|AF361470_8 hypothetical protein [Rhizobium Length = 302 Best-BlastP=> >nrprot 57% Legionella longbeachae] 1386.3
 - Identities = 191/434 (44%), Positives = 267/434 (61%), Gaps = 11/434 (2%) ref[ZP_00030193.1| COG1538: Outer Length = 509 Length = 115 membrane protein [Burkholderia fungorum] Best-BlastP=> >nrprot 58% eguminosarum bv. trifolii] 1387.2
 - Best-BlastP=> >nrprot No Hits found 1388.2
- Best-BlastP=> >nrprot 10% Identities = 77/305 (25%), Positives = 150/305 (49%), Gaps = 30/305 (9%) gbjAAK01145.2| 200 kDa Length = 1421 mmunoreactive glycoprotein [Ehrlichia canis] 139.5
 - Best-BlastP=> >nrprot No Hits found 1390.5
- Best-BlastP=> >nrprot 41% Identities = 31/81 (38%), Positives = 45/81 (55%), Gaps = 2/81 (2%) ref[ZP_00092569.1| COG3293: Transposase Identities = 119/193 (61%), Positives = 151/193 (78%), Gaps = 1/193 (0%) ref|NP_820272.1| proton transporter, Length = 200putative [Coxiella burnetii RSA 493] gb|AAO90786.1| proton transporter, putative [Coxiella burnetii RSA 493] Best-BlastP=> >nrprot 73% 1392.2
 - Length = 249vinelandii] and inactivated derivatives [Azotobacter
- Best-BlastP=> >nrprot 63% Identities = 55/150 (36%), Positives = 95/150 (63%), Gaps = 1/150 (0%) ref|ZP_00031963.1| COG0517: FOG: CBS Length = 150domain [Burkholderia fungorum] 1394.2
- 7120] pir]|AD1916 alcohol dehydrogenase [imported] Nostoc sp. (strain PCC 7120) dbj|BAB72836.1| alcohol dehydrogenase [Nostoc sp. PCC Identities = 204/328 (62%), Positives = 250/328 (76%) ref[NP_484922.1] alcohol dehydrogenase [Nostoc sp. PCC Best-BlastP=> >nrprot 75% Length = 328 1396.3
- Identities = 56/149 (37%), Positives = 86/149 (57%), Gaps = 4/149 (2%) ref[NP_771277.1] blr4637 [Bradyrhizobium Length = 173 aponicum] dbj|BAC49902.1| blr4637 [Bradyrhizobium japonicum USDA 110] Best-BlastP=> >nrprot 52% 1397.2 1398.2
- Identities = 135/479 (28%), Positives = 236/479 (49%), Gaps = 16/479 (3%) refINP_460415.1| putative POT family, Best-BlastP=> >nrprot 56% Identities = 261/666 (39%), Positives = 375/666 (56%), Gaps = 31/666 (4%) ref[NP_901612.1] probable peptidase typhimurium LT2] gb[AAL20374.1] putative POT family peptide transport protein [Salmonella [Chromobacterium violaceum ATCC 12472] gb[AAQ59616.1] probable peptidase [Chromobacterium violaceum ATCC 12472] peptide transport protein [Salmonella Best-BlastP=> >nrprot 47% 1399.3

Identities = 245/484 (50%), Positives = 346/484 (71%), Gaps = 8/484 (1%) refINP_820418.1| NADH dehydrogenase I, N subunit [Coxiella burnetii RSA 493] gb|AAO90932.1| NADH dehydrogenase I, N subunit [Coxiella burnetii RSA 493] Best-BlastP=> >nrprot 71% 1402.2

1400.2

- Identities = 289/499 (57%), Positives = 367/499 (73%), Gaps = 1/499 (0%) ref[NP_820419.1] NADH dehydrogenase , M subunit [Coxiella burnetii RSA 493] gb|AAO90933.1| NADH dehydrogenase I, M subunit [Coxiella burnetii RSA 493] Best-BlastP=> >nrprot 73%
 - Best-BlastP=> >nrprot 98% Identities = 292/296 (98%), Positives = 293/296 (98%) emb|CAB43070.1| DjlA protein [Legionella pneumophila] 1405.2
- Best-BlastP=> >nrprot 97% Identities = 401/419 (95%), Positives = 408/419 (97%) emb|CAB43071.1| 3-Deoxy-D-manno-oct-2-ulosonic acid Length = 419 pneumophila] ransferase [Legionella 1406.2
 - 1407.2 Best-BlastP=> >nrprot No Hits found

ength = 296

- Best-BlastP=> >nrprot 60% Identities = 102/245 (41%), Positives = 157/245 (64%), Gaps = 3/245 (1%) ref[NP_819783.1| competence lipoprotein ComL, putative [Coxiella burnetii RSA 493] gb|AAO90297.1| competence lipoprotein ComL, putative [Coxiella burnetii RSA 493] 1409.3
- Best-BlastP=> >nrprot 25% Identities = 116/527 (22%), Positives = 221/527 (41%), Gaps = 72/527 (13%) pir||T18296 myosin heavy chain -Length = 2139 Entamoeba histolytica gb|AAB48065.1| myosin heavy chain [Entamoeba histolytica] 141.4
 - Best-BlastP=> >nrprot 50% Identities = 94/303 (31%), Positives = 159/303 (52%), Gaps = 8/303 (2%) refINP_931129.1| recombinaison laumondii TTO1] emb|CAE16301.1| recombinaison associated protein Length = 303laumondii TTO11 associated protein [Photorhabdus luminescens subsp. Photorhabdus luminescens subsp. 1410.2
- Best-BlastP=> >nrprot 61% Identities = 251/573 (43%), Positives = 359/573 (62%), Gaps = 17/573 (2%) ref[ZP_00110863.1| hypothetical protein Length = 578Nostoc punctiforme] 1412.5
 - Best-BlastP=> >nrprot 36% Identities = 99/349 (28%), Positives = 168/349 (48%), Gaps = 18/349 (5%) refINP_284204.1| putative acyl-CoA Neisseria meningitidis Lenath = 517igase [Neisseria meningitidis Z2491] pir||D81839 probable acyl-CoA ligase (EC 6.2.1.-) NMA1482 [imported] strain Z2491 serogroup A) emb[CAB84715.1| putative acyl-CoA ligase [Neisseria meningitidis Z2491] 1413.2
 - Best-BlastP=> >nrprot 52% Identities = 24/70 (34%), Positives = 41/70 (58%), Gaps = 2/70 (2%) ref[ZP_00036213.1| hypothetical protein Length = 77. Enterococcus faecium] 1414.1
- Identities = 75/253 (29%), Positives = 121/253 (47%), Gaps = 30/253 (11%) ref|ZP_00012801.1| hypothetical protein Length = 255Rhodopseudomonas palustris] Best-BlastP=> >nrprot 48% 1415.4 1417.1
 - Best-BlastP=> >nrprot 66% Identities = 64/129 (49%), Positives = 88/129 (68%) ref|ZP_00066647.1| hypothetical protein [Microbulbifer Length = 164 degradans 2-40) 1418.3
- Best-BlastP=> >nrprot 71% Identities = 295/592 (49%), Positives = 421/592 (71%), Gaps = 2/592 (0%) ref[NP_820066.1| transporter, putative Coxiella burnetii RSA 493] gb|AAO90580.1| transporter, putative [Coxiella burnetii RSA 493]